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UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new nonprovisional applications under 37 C.F.R. § 1.53(b))

Attorney Docket No. LA0046a
First Inventor or Application Identifier Beth Anne Piper
Title METHOD FOR TREATING DIABETES
Express Mail Label No. EM069782219US

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

1. ☒ * Fee Transmittal Form (e.g., PTO/SB/17)
(Submit an original and a duplicate for fee processing)
2. ☒ Specification [Total Pages 53]
(preferred arrangement set forth below)
 - Descriptive title of the invention
 - Cross References to Related Applications
 - Statement Regarding Fed sponsored R & D
 - Reference to Microfiche Appendix
 - Background of the invention
 - Brief Summary of the invention
 - Brief Description of the Drawings (if filed)
 - Detailed Description
 - Claim(s)
 - Abstract of the Disclosure
3. ☒ Drawing(s) (35 U.S.C. 113) [Total Sheets 10]
4. Oath or Declaration [Total Pages 3]
 - a. ☒ Newly executed (original or copy)
 - b. ☐ Copy from a prior application (37 C.F.R. § 1.63(d))
(for continuation/divisional with Box 16 completed)
 - i. ☐ DELETION OF INVENTOR(S)
Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §§ 1.63(d)(2) and 1.33(b).

***NOTE FOR ITEMS 1 & 13: IN ORDER TO BE ENTITLED TO PAY SMALL ENTITY FEES, A SMALL ENTITY STATEMENT IS REQUIRED (37 C.F.R. § 1.27), EXCEPT IF ONE FILED IN A PRIOR APPLICATION IS RELIED UPON (37 C.F.R. § 1.28).**

ADDRESS TO: Assistant Commissioner for Patents
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5. ☐ Microfiche Computer Program (Appendix)
6. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary)
 - a. ☐ Computer Readable Copy
 - b. ☐ Paper Copy (identical to computer copy)
 - c. ☐ Statement verifying identity of above copies

ACCOMPANYING APPLICATION PARTS

7. ☒ Assignment Papers (cover sheet & document(s))
8. ☐ 37 C.F.R. § 3.73(b) Statement of Power of Attorney (when there is an assignee)
9. ☐ English Translation Document (if applicable)
10. ☐ Information Disclosure Statement (IDS)/PTO-1449 [Copies of IDS Citations]
11. ☐ Preliminary Amendment
12. ☒ Return Receipt Postcard (MPEP 503)
(Should be specifically itemized)
13. ☐ * Small Entity Statement(s) [Statement filed in prior application, Status still proper and desired (PTO/SB/09-12)]
14. ☐ Certified Copy of Priority Document(s) (if foreign priority is claimed)
15. ☐ Other: _____

16. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below and in a preliminary amendment:

☐ Continuation ☐ Divisional ☒ Continuation-in-part (CIP) of prior application No: 09,432,465
Prior application information: Examiner _____ Group / Art Unit: 1646

For CONTINUATION or DIVISIONAL APPS only: The entire disclosure of the prior application, from which an oath or declaration is supplied under Box 4b, is considered a part of the disclosure of the accompanying continuation or divisional application and is hereby incorporated by reference. The incorporation can only be relied upon when a portion has been inadvertently omitted from the submitted application parts.

17. CORRESPONDENCE ADDRESS

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Bristol-Myers Squibb Company

P.O. Box 4000 Princeton, NJ 08543-4000 609 252-4000

Patent Department

Case No.: LA0046a

To the Assistant Commissioner for Patents:
Washington, D.C. 20231

Sir:

Forwarded herewith is a patent application consisting of specification, claims, Declaration, 10 sheet(s) of drawing and Assignment. The title is: METHOD FOR TREATING DIABETES

The inventor(s) is (are): Beth Anne Piper

The filing fee is believed to be as follows:

Basic fee: \$760.00

Additional fees:

Total number of claims in
excess of 20, times \$18.00 (16) 288.00

Number of independent claims
minus 3, times \$78.00 156.00

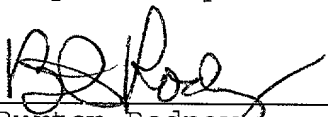
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Respectfully submitted,


Burton Rodney
Attorney

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Date of Deposit: Dec. 14, 1988

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METHOD FOR TREATING DIABETESReference to Other Applications

This is a continuation-in-part of U.S. Application
5 Serial No. 09/432,465, filed November 3, 1999 (Attorney
File LA0046).

Field of the Invention

The present invention relates to a method for
10 treating type 2 diabetes in drug naive patients,
employing a low dose formulation which includes metformin
and glyburide. The low dose formulation has at least
substantially equivalent efficacy in treating type 2
diabetes as compared to formulations containing higher
15 doses of metformin and/or glyburide, but with
substantially reduced side effects.

Background of the Invention

The biguanide antihyperglycemic agent metformin
20 disclosed in U.S. Patent No. 3,174,901 is currently
marketed in the U.S. in the form of its hydrochloride
salt (Glucophage®), Bristol-Myers Squibb Company).

The diagnosis and management of type 2 diabetes
mellitus is rapidly undergoing progressive changes. It
25 is now widely accepted that glycemic control makes a
difference. The goal of diabetes therapy today is to
achieve and maintain as near normal glycemia as possible
to prevent the long-term microvascular and macrovascular
complications of an elevated blood glucose. The
30 diagnosis of diabetes has undergone significant changes
as evidenced by the new ADA diagnostic and classification
guidelines. Oral therapeutic options for the treatment
of type 2 diabetes mellitus, until recently, have been
severely limited. Prior to 1995, sulfonyl ureas had been
35 the mainstay of oral diabetes agents in the United
States. Sulfonyl ureas target one mechanism of
hyperglycemia by augmenting insulin secretion from the

beta cell. Since 1995, three new classes of agents have been added to the anti-diabetes armamentarium for the management of hyperglycemia. Metformin, a biguanide, targets additional mechanisms of hyperglycemia by
5 inhibiting hepatic glucose production and enhancing peripheral glucose uptake and thereby reduce insulin resistance; thiazolidinediones such as troglitazone, rosiglitazone and pioglitazone decrease peripheral
10 insulin resistance; and alpha-glucosidase inhibitors such as acarbose and miglitol help control postprandial glucose excursion by delaying absorption of dietary carbohydrate. These agents are all indicated as monotherapy and some are indicated for use in combination therapy, generally, after monotherapy has been found to
15 be inadequate.

In 1995, metformin was added to sulfonyl urea therapy in patients who had not achieved glycemic control with sulfonyl urea monotherapy and the two agents were found to have a remarkable effect on glycemic control or
20 lowering of hemoglobin-A1c. The different mechanisms of action in targeting hyperglycemia are complimentary and make combination use attractive and a rational course of action. Prescription data reveals approximately 60% of metformin use is in combination with a sulfonyl urea.

25 Examples of combinations of metformin and the sulfonyl urea glyburide (also referred to as glibenclamide) are disclosed in the following references.

(1) WO 97/17975 published May 22, 1997, (Barelli et al, Istituto Gentili S.P.A.) and U.S. Patent No.
30 5,922,769 to Barelli et al (hereinafter Barelli et al) discloses a combination of glibenclamide and metformin in a 1:100 weight ratio, so as to allow a daily dosage of 15 mg glibenclamide and 1500 mg metformin, used for the onset of diabetes to the most severe cases, particular in
35 cases of secondary failure to a combination of glibenclamide-metformin HCl in a weight ratio higher than 1:100.

(2) Vigneri et al, Treatment of NIDDM Patients with Secondary Failure to Glyburide: Comparison of the Addition of Either Metformin or Bed-Time NPH Insulin to Glyburide, *Diabete & Metabolisme*, 1991, 17, 232-234, disclose use of a combination of 1.5 g/day metformin and 15 mg/day glyburide to treat NIDDM patients with secondary failure to 15 mg/day glyburide.

(3) Higginbotham et al, Double-Blind Trial of Metformin in the Therapy of Non-Ketotic Diabetes, *The Medical Journal of Australia*, August 11, 1979, 154-156, discloses treatment of diabetic patients, who were already receiving from 10 mg to 20 mg per day of glibenclamide, with 500 mg metformin twice a day. Higginbotham et al conclude "that in selected diabetics whose condition is inadequately controlled with sulphonylurea therapy, significant improvement in diabetic control can be obtained by the addition of metformin in a low dose of 500 mg twice a day."

(4) U.S. application Serial No. 09/353,141, filed July 14, 1999 (based on European application No. 98401781.4, filed July 15, 1998) discloses formulations containing metformin and glyburide where the glyburide is of a particular particle size as described hereinafter.

References which disclose combinations of metformin and glipizide include the following:

- (1) Combination of glipizide/metformin treatment reduces low density lipoprotein binding to arterial proteoglycans in DIDDM, Edwards et al, *Diabetes*, (46, Suppl. 1, 45A, 1997).
- (2) Combination of glipizide/metformin normalizes glucose and improves insulin sensitivity in hyperinsulinemia moderately well controlled. Cefalu et al, *Diabetes*, (45, Suppl. 2, 201A, 1996).
- (3) Effects of combination of glipizide/metformin treatment on oxidizability of LDL in NIDDM, Crouse et al, *Circulation*, (94, No. 8, Suppl., I508, 1996).

(4) Insulin sensitivity is improved after glipizide monotherapy and combination with metformin, Cefalu et al, Diabetologia, (39, Suppl. 1, A231, 1996).

5 (5) Combined Metformin - Sulfonyl urea Treatment of Patients with NIDDM in Fair to Poor Glycemic Control, Reaven et al, J. Clin. Endocrinol. Metab. (74, No. 5, 1020-26, 1992).

10 (6) Combination of Glipizide/Metformin Treatment in NIDDM, Hollenbeck et al, Diabetes, (39, Suppl. 1, 108A, 1990).

(7) Oral Antidiabetic Combination Therapy with Sulfonyl ureas and Metformin, Haupt et al, Med. Welt. (40, No. 5, 118-23, 1989).

15 (8) Variation of the lipemic pattern in diabetic subjects after treatment with a combination of glipizide and metformin, Ferlito et al, PROGR. MED. (Roma) 31/6 (289-301) 1975.

(9) Results with a combination of glipizide and dimethylbiguanide in 40 cases of diabetes, Parodi et al, 20 GAZZ. MED. ITAL. 132/5 (226-235) 1973.

Other combinations of metformin and another antidiabetic agent are disclosed in the following references.

25 (1) U.S. Patent No. 5,631,224 to Efendic et al discloses a combination of metformin with GLP-1(7-36) amide or GLP-1(7-37) or a fragment thereof.

(2) WO 98/57634 (SKB) discloses a method for treating diabetes employing a combination of a thiazolidinedione and metformin. The thiazolidinedione 30 may be troglitazone, ciglitazone, pioglitazone or englitazone, and may be employed in dosages of 2 to 12 mg per day while the metformin may be employed in daily dosages "of up to 3000 mg per day, in unit doses of 500 mg (for example, 2 to 3 times per day) or 850 mg (2 times 35 per day), one example of a dosage for metformin is 500 mg building to 5 times per day."

(3) U.S. Patent No. 5,965,584 (Takeda) discloses a combination of a thiazolidenedione insulin sensitivity enhancer (such as pioglitazone) and metformin.

None of the above references suggests employing
5 diabetic combinations containing metformin for first line treatment of drug naive patients.

Several fixed combinations of metformin and glyburide (glibenclamide) are presently being marketed outside the U.S. These include (1) combinations of 400
10 mg metformin/2.5 mg glibenclamide (Boehringer's Bi-Euglucon in Argentina, and Bi-Euglicon M in Italy; Guidotti/Menarini's Glibomet in the Dominican Republic and Italy; HMR's Normell in Greece and Hoechst's Suguan-M in Italy; Sun Pharma's Glucored in India; Monsanto's
15 (Searle's) Benclamet in India; Guidotti's Glibomet in Liban; Berlin Chemie/Menarini's Glibomet in the Slovak Rep., and Roche's Bi-Euglucon in Uruguay); (2) combinations of 500 mg metformin/5 mg glibenclamide (Sun Pharma's Glucored in India; Monsanto's (Searle's)
20 Benclamet in India, USV's Duotrol in India; and Lakeside's (Roche) Bi-Euglucon M5 in Mexico); (3) combinations of 500 mg metformin/2.5 mg glibenclamide (Molteni's Glucomide in Italy, Lakeside's (Roche) Bi-Euglucon M in Mexico and Szabo's Dublex in Uruguay); and
25 (4) 1 g metformin/5mg glibenclamide (Silanes Sil-Norboral in Mexico).

The labelling for Glucophage® (Bristol-Myers Squibb's metformin), in the Physicians' Desk Reference 1999, under "Indications and Use", indicates that
30 Glucophage may be used concomitantly with a sulfonylurea. It is further indicated under "Dosage and Administration" "Concomitant Glucophage and Oral Sulfonylurea Therapy" that "If patients have not responded to four weeks of the maximum dose of Glucophage monotherapy, consideration
35 should be given to gradual addition of an oral sulfonylurea while continuing Glucophage at the maximum dose.... With concomitant Glucophase and sulfonylurea

therapy, the desired control of blood glucose may be obtained by adjusting the dose of each drug. However, attempts should be made to identify the maximum effective dose of each drug to achieve this goal." The recommended dosing schedule for Glucophage is a starting dose of 500 mg twice a day or 850 mg once a day with dosage increases in increments of 500 mg weekly or 850 mg every 2 weeks up to a total of 2000 mg per day.

Package inserts for Bi-Euglucon M and Suguan M in Italy (400 mg metformin/2.5 mg glibenclamide) indicate that these drug combinations are used in cases of primary or secondary resistance to sulfonyl ureas [that is as second or third line therapy] and that a dosage of $\frac{1}{2}$ tablet per day increasing $\frac{1}{2}$ tablet at a time according to glycemic variations up to 4 tablets per day are employed.

Package inserts for Glibomet (400 mg metformin/2.5 mg glibenclamide) and Glucomide (500 mg metformin/2.5 mg glibenclamide) in Italy indicate that these drug combinations are used for treating type 2 diabetes which is non-controllable or cannot be controlled with only diet or with diet and sulfonyl urea [that is as first line therapy or second line therapy].

The package insert for Glibomet in Italy indicates a daily dosage of 2 tablets, that is 800 mg metformin and 5 mg glibenclamide, up to 2 grams metformin. The package insert for Glucomide in Italy indicates a daily dosage of 2 capsules, that is 1000 mg metformin up to 2 grams metformin, and 5 mg glibenclamide.

Description of the Invention

In accordance with the present invention, a method is provided for treating diabetes, especially type 2 diabetes, in a drug naive human patient, which includes the step of administering to a drug naive human patient in need of treatment, as first line therapy, a therapeutically effective low dose pharmaceutical formulation which includes a combination of metformin and

glyburide. The above combination will preferably provide at least substantially equivalent efficacy in treating diabetes in drug naive patients as do combinations of metformin and glyburide employed in higher dosages, such as prescribed in generally accepted medical practice for first line therapy treating diabetes, but with substantially reduced side effects.

In one aspect of the method of the invention, the daily dosage of metformin administered will be less than 800 mg.

It is to be understood that the low dose formulation employed in the method of the invention will include a starting "low dose" of the active antidiabetes drug components, that is a lower starting dosage than the starting dosage for such drug prescribed in generally accepted medical practice in first line therapy of treating diabetes. Thus, the above low dose pharmaceutical formulation will include a low dose of metformin and a low dose of glyburide as defined hereinafter.

In accordance with the present invention, efficacy in first line therapy in treating diabetes in drug naive patients is achieved employing the low dose pharmaceutical formulation wherein the starting daily dosage of metformin is as low as about one-fifth of the starting daily dosage of metformin employed in generally accepted medical practice for first line therapy for treating diabetes (that is a starting daily dosage of as low as 160 mg metformin per day), up to a daily maintenance dosage of metformin employed in generally accepted medical practice for first line or second line therapy for treating diabetes (that is up to 2000 mg metformin per day). Preferably, the maximum daily maintenance dosage of metformin will be about two-thirds of the daily maintenance dosage of metformin employed in generally accepted medical practice for first line therapy for treating diabetes.

In carrying out the method of the invention, the starting daily dosage of metformin will be as low as from about 25% up to about 60% of the starting daily dosage of metformin employed in generally accepted medical practice for first line therapy for treating diabetes (that is a starting daily dosage from 160 to 500 mg metformin, preferably 250 or 500 mg metformin). Where necessary, the starting dosage may be titrated up to a daily maintenance dosage of from about 40 to about 100%, preferably from about 40 to about 60% of the maintenance dosage employed in generally accepted medical practice for first line therapy for treating diabetes (that is a daily maintenance dosage from 320 to 2000 mg, preferably from 320 to 1200 mg).

In the method of the invention, the low dose pharmaceutical formulation will preferably be employed in first line therapy in a daily dosage to provide less than about 800 mg metformin per day, preferably no more than about 750 mg metformin per day, more preferably no more than about 600 mg metformin per day, and a starting dosage of from about 160 to about 500 mg per day, preferably 250 mg per day or 500 mg per day, in single or divided doses of one to four tablets daily.

The glyburide is employed in a starting daily dosage as low as about one-fifth of the starting daily dosage of glyburide employed in generally accepted medical practice for first line or second line therapy for treating diabetes (that is a minimum starting daily dosage as low as 0.5 mg). Where necessary, the starting dosage of glyburide may be titrated up to a daily maintenance dosage of glyburide employed in generally accepted medical practice for first or second line therapy for treating diabetes (that is up to a maximum of 15 mg glyburide per day). Preferably, the maximum daily dosage of glyburide will be about two-thirds of the daily maintenance dosage of glyburide employed in generally accepted medical practice for first line therapy for

treating diabetes (that is up to a maximum of 2.5 to 10 mg glyburide per day).

5 The glyburide will preferably be employed in a starting daily dosage as low as about 20% up to about 60% of the starting daily dosage of glyburide employed in generally accepted medical practice for first line therapy for treating diabetes (that is a minimum starting dosage as low as 0.5 mg to 3.5 mg, more preferably 1.25 mg or 2.5 mg). The glyburide may be titrated up to a
10 daily maintenance dosage of about 40 to about 100%, preferably from about 40 to about 60% of the daily maintenance dosage of glyburide employed in generally accepted medical practice for first line therapy for treating diabetes (that is a maximum daily dosage of 2 to
15 15 mg preferably a maximum daily dosage of 2.5 to 10 mg).

The above daily dosage of glyburide may be employed in single or divided doses of one to four tablets daily.

The metformin and glyburide may be formulated in a single tablet which may be employed in single or divided
20 doses of one to four times daily.

The term "low dose combination", "low dose formulation" or "low dose pharmaceutical formulation" as employed herein, in a most preferred formulation, refers to a formulation which includes as a starting daily
25 dosage 250 mg metformin, and 1.25 mg glyburide, or 500 mg metformin and 2.5 mg glyburide.

Until now, combinations of metformin and glyburide, have normally been used with few exceptions, as second line therapy in treating type 2 diabetes. Generally
30 accepted medical practice daily dosages for such second line therapy employing fixed combinations of metformin and glyburide range from 3 to 4 tablets containing 400 to 500 mg metformin and 2 to 2.5 mg glyburide, or about 1200 to 2000 mg metformin and 6 to 10 mg glyburide, daily.

35 As indicated above with respect to Glibomet and Glucoside (fixed combinations of metformin and glyburide) marketed in Italy, these combinations may be employed as

first line therapy (drug naive patients) in a daily dosage of 800 to 1000 mg up to 2 grams metformin and 5 mg glibenclamide (glyburide).

5 The above dosages may be included within the term dosages as prescribed in generally accepted medical practice for first line therapy or second line therapy in treating diabetes. In some refractory cases of diabetes, up to 15 mg glyburide may be indicated.

10 As indicated above with respect to Boehringer's Bi-Euglucon M and Hoechst's Suguan M (fixed combinations of metformin and glibenclamide) marketed in Italy, these combinations are employed as second line therapy in a daily dosage starting at $\frac{1}{2}$ tablet, that is, 200 mg metformin and 1.25 mg glibenclamide. The initial or
15 starting low doses are employed to determine if the patient can tolerate the drugs. Furthermore, there apparently is no known clinical first line therapy study available which supports use of these starting doses. These starting doses are gradually titrated upwardly $\frac{1}{2}$
20 tablet at a time up to 4 tablets per day until an efficacious dosage is achieved. Thus, the initial or starting daily dosage of $\frac{1}{2}$ tablet or 200 mg metformin and 1.25 mg glibenclamide is not considered herein as "dosages as prescribed in generally accepted medical
25 practice for treating diabetes".

Surprisingly, it has been found that use of the combination of metformin and glyburide in accordance with the present invention affords the following benefits. The low dose metformin is an insulin sensitizer and
30 decreases insulin resistance at the liver, muscle and pancreas. The low dose metformin-glyburide combination acts on the pancreas as a glucose sensitizer; it decreases glucose toxicity at the pancreas and improves function of the pancreas.

35 In addition, in accordance with the present invention, a method is provided for treating diabetes, especially type 2 diabetes, in a drug naive human

patient, which includes the step of administering to a drug naive human patient in need of treatment, as first line therapy, a starting low dose pharmaceutical formulation which includes a combination of metformin and glyburide. The starting low dose combination preferably provides at least substantially equivalent efficacy in treating diabetes in drug naive patients as do combinations of metformin and glyburide employed in dosages (including starting dosages) prescribed in generally accepted medical practice for first line therapy treating diabetes, but with substantially reduced side effects.

The starting daily dosage of metformin is as low as 20% of the starting daily dosage of metformin employed in generally accepted medical practice for first line therapy for treating diabetes, preferably a starting daily dosage from about 160 to about 500 mg, more preferably a starting daily dosage of 250 mg or 500 mg.

The starting daily dosage of glyburide is as low as 20% of the starting daily dosage of glyburide employed in generally accepted medical practice for first line therapy for treating diabetes, preferably a starting daily dosage from about 0.625 to about 5 mg, more preferably a starting daily dosage of 1.25 mg or 2.5 mg.

In addition, in accordance with the present invention, a method is provided for decreasing fasting plasma glucose, decreasing insulin resistance, decreasing hemoglobin A1c, increasing post-prandial insulin and/or decreasing post-prandial glucose excursion in a human diabetic patient, which includes the step of administering as first line therapy to a drug naive human patient a low dose pharmaceutical formulation which includes a combination of metformin and glyburide.

In carrying out the method of the invention employing the preferred starting low dose pharmaceutical formulation containing metformin and glyburide, to treat drug naive patients for diabetes, the efficacy in

treating drug naive patients is at least substantially equivalent and incidence of side effects (gastrointestinal side effects and hypoglycemia) is surprisingly significantly and substantially reduced as compared to patients on higher daily dosages of metformin and glyburide (that is in starting dosages prescribed in generally accepted medical practice for treating diabetes). Thus, while efficacy in treating drug naive patients as measured by decrease in hemoglobin A_{1c} (HbA_{1c}) from baseline over time, decrease in fasting plasma glucose (FPG), increase in post-prandial insulin levels, and decrease in post-prandial glucose (PPG) excursion, are essentially substantially equivalent in the above-described patients when employing the low dose pharmaceutical formulation employed herein and substantially higher daily dosages, incidence of hypoglycemia and gastrointestinal side effects in drug naive patients treated with substantially higher daily dosages are substantially greater than in patients treated with the low dose pharmaceutical formulation.

Most preferred dosages for use herein are 250 mg metformin/1.25 mg glyburide, and 500 mg metformin/2.5 mg glyburide.

The low dose pharmaceutical formulation of metformin and glyburide is employed as initial therapy that is as an adjunct to diet and exercise to improve glycemic control in patients with type 2 diabetes mellitus.

The ADA recommends a treatment goal of HbA_{1c} < 7% (ADA. *Diabetes Care* 21 [Suppl. 1]: S23 - S31, 1998) in order to reduce the risk of complications of type 2 diabetes mellitus, including coronary heart disease and microvascular complications.

Dosage of the preferred metformin-glyburide combination must be individualized on the basis of both effectiveness and tolerance. It is preferably given with meals and should be started at a low dose, with gradual

dose escalation. Ideally, the response to therapy should be evaluated using HbA_{1c} (glycosylated hemoglobin) which is a better indicator of long-term glycemic control than FPG alone. The therapeutic goal in all patients with
5 type 2 diabetes mellitus should be to improve glycemic control, including FPG, postprandial glucose and HbA_{1c} levels, to normal or as near normal as possible. Patients should be titrated to achieve the ADA goal of HbA_{1c} < 7% following the dosing recommendations up to the
10 maximum recommended dose. (ADA. *Diabetes Care* 21 [Suppl. 1]: S23 - S32, 1998).

As initial therapy, the most preferred starting dose of the metformin-glyburide combination is 250/1.25 mg once a day, given with a meal. For patients with a
15 baseline HbA_{1c} > 9% or a fasting glucose > 200 mg/dL, a recommended starting dose of 250/1.25 mg twice daily with the morning and evening meal may be preferred. Dosage increases should preferably be made in increments of 250/1.25 mg, every 2 weeks, up to the minimum effective
20 dose necessary to achieve adequate glycemic control. For those patients requiring additional glycemic control, the 250 mg/1.25 mg dosage may be switched to 500/2.5 mg.

The preferred low dose metformin-glyburide
25 formulation are set out below.

Product identity	Amount of ingredient, mg per tablet
	250/1.25 or 500/2.5 or 500/5.0
Ingredient	
Metformin hydrochloride	250.0 or 500.0
Glyburide	1.25 or 2.5 or 5
Croscarmellose sodium	3.0-15.0
Microcrystalline cellulose	15.0-60.0
Polyvinyl pyrrolidone	3.0-20
Magnesium stearate	0.3-7.5
Film coat*	4.5-12.0

*a commercially available film coat composition is used, such as Opadry (Colorcon, UK).

- 5 The especially preferred low dose metformin-glyburide formulations are as follows:

Product identity	Amount of ingredient, mg per tablet		
	250/1.25	500/2.5	500/5.0
Ingredient			
Metformin hydrochloride*	251.25	502.50	502.50
Glyburide	1.25	2.5	5.0
Croscarmellose sodium	7.0	14	14
Microcrystalline cellulose	28.25	56.50	54.0
Polyvinyl pyrrolidone	10.0	20	20
Magnesium stearate	2.25	4.50	4.50
Film coat**	6	12.0	12.0

*contains 99.5% metformin HCl and 0.5% Mg stearate (w/w)

**a commercially available film coat composition is used, such as Opadry (Colorcon, UK).

10

- 15 The low dose pharmaceutical formulation containing the metformin-glyburide combination, will preferably be formulated according to the teachings disclosed in U.S. application Serial No. 09/353,141, filed July 14, 1999, which claims priority from European application No. 98401781.4 filed July 15, 1998, which U.S. application is incorporated herein by reference.

The preferred low dose pharmaceutical formulation employed in the method of the invention in the form of a solid oral form such as a tablet will contain the metformin-glyburide combination as disclosed in U.S.

5 Application Serial No. 09/353,141, filed July 14, 1999, and as such will include glyburide which has a glyburide bioavailability comparable to the glyburide availability obtained with a separate administration of metformin and glyburide. This is accomplished by employing glyburide
10 in a predetermined particle size distribution. Thus, the metformin-glyburide formulation will contain a combination of metformin and glyburide in which the size of the glyburide is such that at most 10% of the particles are less than 2 μm and at most 10% of the
15 particles are greater than 60 μm . Preferably, the size of the glyburide is such that at most 10% of the particles are less than 3 μm and at most 10% of the particles are greater than 40 μm . This specific size range of glyburide may be obtained by sieving or air jet
20 milling.

In a second embodiment, the low dose solid oral dosage form will contain a combination of metformin and glyburide in which the size of glyburide is such that at most 25% of the particles are less than 11 μm and at most
25 25% of the particles are greater than 46 μm .

Preferably, 50% of particles are less than 23 μm .

Most preferred are a combination of metformin and glyburide, where the glyburide has a particle size distribution of about 25% undersize value not more than 6
30 μm , about 50% undersize value 7 to 10 μm and about 75% undersize value not more than 23 μm .

Detailed Description of the Invention

The term "diabetes" as employed herein, refers to
35 type 2 (or Type II) diabetes or non-insulin dependent diabetes mellitus (NIDDM).

The term "metformin" as employed herein refers to metformin or a pharmaceutically acceptable salt thereof such as the hydrochloride salt, the metformin (2:1) fumarate salt, and the metformin (2:1) succinate salt as
5 disclosed in U.S. application Serial No. 09/262,526 filed March 4, 1999, the hydrobromide salt, the p-chlorophenoxy acetate or the embonate, and other known metformin salts of mono and dibasic carboxylic acids including those disclosed in U.S. Patent No. 3,174,901, all of which
10 salts are collectively referred to as metformin. It is preferred that the metformin employed herein be the metformin hydrochloride salt, namely, that marketed as Glucophage® (trademark of Bristol-Myers Squibb Company).

The term "substantially reduced side effects" as
15 employed herein refers to reduced incidence of hypoglycemia and gastrointestinal adverse events including diarrhea, nausea/vomiting and/or abdominal pain, occurring with use of the low dose pharmaceutical formulation in drug naive patients as compared to
20 patients on the same active components in the pharmaceutical formulation of the invention but at higher dosages.

The term "at least substantially equivalent efficacy" in treating type 2 diabetes as employed herein
25 refers to the effectiveness of the low dose pharmaceutical formulation in treating drug naive patients to reduce and/or maintain hemoglobin A_{1c} (glycosylated hemoglobin) at 7% or less, to decrease insulin resistance (by increasing post-prandial insulin level) and/or to decrease post-prandial glucose (PPG)
30 excursion, as compared to patients treated with the same active components in the pharmaceutical formulation of the invention but at higher dosages.

The term "post-prandial excursion" as employed
35 herein refers to the difference between post-prandial glucose (PPG) and fasting plasma glucose (FPG).

The low dose pharmaceutical formulation containing metformin in combination with glyburide may be administered orally in the same dosage form or in separate oral dosage forms or by injection.

5 It is believed that the use of metformin in combination with glyburide produces antihyperglycemic results greater than that possible from each of these medicaments alone and greater than the combined additive anti-hyperglycemic effects produced by these medicaments.

10 Metformin will be employed in a weight ratio to the glyburide in the range from about 1000:1 to about 10:1, preferably from about 400:1 to about 50:1, more preferably from about 200:1 to about 100:1.

15 In carrying out method of the present invention, a low dose pharmaceutical formulation or composition will be employed containing metformin and glyburide in association with a pharmaceutical vehicle or diluent. The low dose pharmaceutical formulation can be formulated employing conventional solid or liquid vehicles or
20 diluents and pharmaceutical additives of a type appropriate to the mode of desired administration. The low dose pharmaceutical formulation can be administered to mammalian species including humans, monkeys, dogs, etc., by an oral route, for example, in the form of
25 tablets, capsules, granules or powders, or it can be administered by a parenteral route in the form of injectable preparations. The dose for drug naive patients is as described above, which can be administered in a single dose or divided doses, from 1-4 times per
30 day.

 The above dosage forms may also include the necessary physiologically acceptable carrier material, excipient, lubricant, buffer, antibacterial, bulking agent (such as mannitol), anti-oxidants (ascorbic acid or
35 sodium bisulfite) or the like.

 The dose administered must be carefully adjusted according to the age, weight, and condition of the

patient, as well as the route of administration, dosage form and regimen, and the desired result.

The combination of the metformin or salt thereof and glyburide may be formulated separately or, where
5 possible, in a single formulation employing conventional formulation procedures.

The various formulations of the invention may optionally include one or more fillers or excipients in an amount within the range of from about 0 to about 90%
10 by weight and preferably from about 1 to about 80% by weight such as lactose, sugar, corn starch, modified corn starch, mannitol, sorbitol, inorganic salts such as calcium carbonate and/or cellulose derivatives such as wood cellulose and microcrystalline cellulose.

One or more binders may be present in addition to or in lieu of the fillers in an amount within the range of from about 0 to about 35% and preferably from about 0.5 to about 30% by weight of the composition. Examples of such binders which are suitable for use herein include
20 polyvinylpyrrolidone (molecular weight ranging from about 5000 to about 80,000 and preferably about 40,000), lactose, starches such as corn starch, modified corn starch, sugars, gum acacia and the like as well as a wax binder in finely powdered form (less than 500 microns)
25 such as carnauba wax, paraffin, spermaceti, polyethylenes or microcrystalline wax.

Where the composition is to be in the form of a tablet, it will include one or more tableting lubricants in an amount within the range of from about 0.2 to about
30 8% and preferably from about 0.5 to about 2% by weight of the composition, such as magnesium stearate, stearic acid, palmitic acid, calcium stearate, talc, carnauba wax and the like. Other conventional ingredients which may optionally be present include preservatives, stabilizers,
35 anti-adherents or silica flow conditioners or glidants, such as Syloid brand silicon dioxide as well as FD&C colors.

Tablets may also include a coating layer which may comprise from 0 to about 15% by weight of the tablet composition. The coating layer which is applied over the outer solid phase containing particles of inner solid phase embedded therein may comprise any conventional coating formulations and will include one or more film-formers or binders, such as a hydrophilic polymer like hydroxypropylmethylcellulose, and/or a hydrophobic polymer like methacrylic acid esters neutral polymer, ethyl cellulose, cellulose acetate, polyvinyl alcohol-maleic anhydride copolymers, β -pinene polymers, glyceryl esters of wood resins and the like and one or more plasticizers, such as triethyl citrate, diethyl phthalate, propylene glycol, glycerin, butyl phthalate, castor oil and the like. Both core tablets as well as coating formulations may contain aluminum lakes to provide color.

The film formers are applied from a solvent system containing one or more solvents including water, alcohols like methyl alcohol, ethyl alcohol or isopropyl alcohol, ketones like acetone, or ethylmethyl ketone, chlorinated hydrocarbons like methylene chloride, dichloroethane, and 1,1,1-trichloroethane.

Where a color is employed, the color will be applied together with the film former, plasticizer and solvent compositions.

The finished dosage form is either a compressed tablet or a hard gelatin capsule, preferably a tablet. The tablet may be optionally film coated. The total amount of drug per dosage unit would be such as to offer a dosage form of convenient size for patients.

The low dose pharmaceutical formulation in the form of a tablet may be obtained by a process as disclosed in U.S. Application Serial No. 09/353,141, filed July 14, 1999, which includes the steps of

- a) forming granules by wet granulation of a mixture of metformin and glyburide

- b) blending the granules with a tableting aid and diluent, and
- c) tableting the blend thus obtained into tablets.

5 The mixture used for forming the granules includes a granulating binder. The granulating binder is preferably a polyvinylpyrrolidone such as, for example, a polyvinylpyrrolidone having a molecular weight of 45,000. The polyvinylpyrrolidone may be used in a proportion of 2
10 to 4% by weight with respect to the final tablet.

 After the granulating step, the granules may be sieved and dried.

 The granules are then blended with a diluent and tableting aid. The diluent may be a conventional filler
15 usually used for making tablets, such as microcrystalline cellulose. The tableting aid may be a conventional material, such as magnesium stearate.

 The tablets thus obtained may then optionally be coated with a hydrophilic cellulose polymer and talc.
20 The hydrophilic cellulose polymer is preferably 2-hydroxypropyl methylcellulose.

Description of the Figures

 Figures 1 and 2 are bar graphs which depict change
25 in hemoglobin Alc (HbAlc) by number of tablets of fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

 Figures 3, 4 and 5 are bar graphs which depict
30 change in HbAlc over time of fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

 Figure 6 is a bar graph which depicts change in
35 fasting plasma glucose (FPG) by number of tablets of fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

Figure 7 is a bar graph which depicts baseline and post-prandial insulin levels of fixed combinations of metformin/glyburide in first line therapy versus monotherapy with glyburide and metformin.

5 Figures 8A and 8B are bar graphs which depict change in PPG excursion at baseline and after 20 weeks of fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

10 Figure 9 is a bar graph which depicts hypoglycemic symptoms in subjects on fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

15 Figure 10 is a bar graph which depicts frequency of gastrointestinal adverse effects in subjects on fixed combinations of metformin/glyburide used in first line therapy versus monotherapy with each of glyburide and metformin.

20 The following Examples represent preferred embodiments of the invention.

Examples 1 to 3

Tablets containing metformin/glyburide combinations were prepared as described below.

25

Composition of Metformin Hydrochloride-Glyburide Tablets
250mg/1.25mg, 500mg/2.5mg and 500mg/5mg

INGREDIENT	Example 1	Example 2	Example 3
	QUANTITY PER TABLET (mg)		
	250mg/1.25mg	500mg/2.5mg	500mg/5mg
Metformin Hydrochloride*	251.25	502.50	502.50
Glyburide	1.25	2.5	5
Croscarmellose Sodium	7.00	14.0	14.0
Povidone	10.00	20.0	20.0
Microcrystalline Cellulose	28.25	56.5	54.0
Magnesium Stearate	2.25	4.5	4.5
Film Coat**	6	12	12

*contains 99.5% metformin HCl and 0.5% Mg stearate (w/w)

**HPMC based film coat used.

5

The metformin hydrochloride-glyburide tablet products, 250 mg/1.25 mg and 500 mg/2.5 mg and 500 mg/5 mg, were compressed from the same granulation. The lower strength tablet was compressed at half the weight of the metformin hydrochloride-glyburide 500 mg/2.5 mg tablet. Tablets manufactured for clinical use were film-coated with a hydroxypropylmethylcellulose (HPMC) film coat. The film coat was non-functional and was applied for aesthetic purposes. The film coat applied to the clinical product was clear.

The manufacturing process for clinical products proceeded as follows:

Croscarmellose sodium and glyburide were dispersed together followed by blending with the metformin hydrochloride/magnesium stearate (99.5%:0.5% w/w) in a high shear mixer. The resultant dry mix was granulated in a high shear mixer with an aqueous povidone solution and dried in a fluid bed dryer at approximately 60°C to achieve a specified moisture content, determined by loss on drying. The dried granulation was reduced with a screening mill and mixed with the microcrystalline cellulose using a tumble mixer. Magnesium stearate was

incorporated as a lubricant using a tumble mixer to produce the final compression blend.

5 The resultant blend was compressed into tablets, to a target weight that was adjusted based on in-process moisture content determinations, on a suitable tablet press. The theoretical tablet weight (based on formula composition with no adjustment for moisture content) was 300 mg for the 250 mg/1.25 mg strength and 600 mg for the 500/2.5 mg strength products.

10 The tablets were film-coated in a perforated coating pan with an appropriate aqueous non-functional HPMC based film coating system until the required amount of film coat had been applied. The typical level of film coat applied to the tablets was 2% w/w.

15 *In vivo* evaluations of prototype combination tablet formulations identified the particle size distribution targeted for use in the clinical program to achieve comparable bioavailability to Micronase from the combination product. The particle size distribution of
20 any glyburide lot was described by three cumulative size criteria: 25% undersize, 50% undersize (also known as the mass median particle size, MMPS) and 75% undersize values. The clinical program involved a total of six glyburide drug substance lots with the 25% undersize
25 value ranging between 4-7 μ m, the 50% undersize value ranging between 8-14 μ m and the 75% undersize value ranging between 17-26 μ m. All six lots of glyburide were synthesized by the same vendor, Profarmaco, with four of them being micronised by Profarmaco. The particle size
30 distributions of the four lots produced are detailed in the following table.

Particle Size Data for Glyburide Drug Substance
Batches Used In Clinical Program

Batch Number	Particle Size ^A (units are equivalent sphere diameters in μm)		
	25% Undersize	50% Undersize	75% Undersize
1	5	9	21
2	5	9	21
3	4	8	18
4	5	9	18

5 ^AParticle size measured by laser light scattering, method reference #CRM 8532 (#SM 248533)

10 The proposed particle size specification included the three cumulative size criteria described above with a range for acceptable mass median particle size (50% undersize) and an upper limit for the lower quartile (25% undersize), and the upper quartile (75% undersize). The particle size specification established for glyburide had been based on the particle size of glyburide used in

15 bioavailability studies, the experience of various clinical lots, the closely matching nature of the size distributions of commercially produced glyburide and the particle size method precision. The particle size criteria described below assured reproducibility of

20 glyburide dissolution and bioavailability from metformin hydrochloride-glyburide tablets.

25 25% undersize value not more than 6 μm
50% undersize value 7-10 μm
75% undersize value not more than 23 μm

Example 4**A. SUMMARY OF 5 CLINICAL PROTOCOLS****(1) Purpose**

5 The following study was conducted to compare
glycemic control of 2 dosage strengths of a fixed
combination metformin/glyburide product (described in
Examples 1 and 2) versus placebo in drug naive patients
with type 2 diabetes mellitus who have had inadequate
10 glycemic control with diet and exercise. The dosage
strengths of fixed combination product evaluated included
metformin 250 mg with glyburide 1.25 mg, and metformin
500 mg with glyburide 2.5 mg. Glycemic control was
assessed using Hemoglobin A1c (HbA_{1c}), the gold-standard
15 measure of long-term glycemic control. Mean change from
baseline in HbA_{1c} following a 20 week treatment period (4
weeks stable once daily dose, 4 week titration and 12
weeks stable dose) were compared. The treatment phase
continued for an additional 12 weeks to assess durability
20 of efficacy.

Contribution of the individual components of the
fixed combination product were assessed by comparison of
short term glycemic parameters of the combination product
and monotherapy arms after 4 weeks of stable once daily
25 dosing. Glycemic control was achieved with similar
incidence of hypoglycemia with the fixed dose
combinations as compared with sulfonyl urea alone or
trends towards decreased gastrointestinal side effects as
compared with metformin alone. Glycemic control was
30 achieved with trends toward decreased adverse events as
compared with either agent alone. Trends in
hypoglycemia, gastrointestinal symptoms and lactate
levels were assessed.

35 (2) Study sites and Subject Population

Eligible subjects were drug naive or have had no
oral antihyperglycemic therapy for the 2 months prior to

screening. Approximately 100 study sites located in the USA were recruited up to a maximum of approximately 800 subjects. Eligible subjects included both males and females between 30 and 78 years of age with established type 2 diabetes mellitus, history of impaired glucose tolerance or impaired fasting glucose who have inadequate glycemic control with diet and exercise.

(3) Study Design and Duration

This study was a 34 week, multicenter, randomized, placebo-controlled, double-blind, parallel study with an optional long-term, open-label treatment phase.

(4) Outcome Measures

Analysis of outcome measures for Periods B and C was performed after all data was made available from the 32 week randomized treatment period.

The primary outcome variable for efficacy was the change from baseline in HbA_{1c} of the two combination therapies relative to placebo following 20 weeks of randomized treatment.

Secondary outcomes included the following:

- Incidence of adverse events, particularly hypoglycemia and gastrointestinal side effects, was compared among treatment arms after 20 and 32 weeks of randomized therapy.
- The number and proportion of subjects achieving a therapeutic glucose response were assessed among treatment arms following 20 and 32 weeks of randomized therapy.
- The reduction in fasting and 2-hour postprandial glucose and insulin were assessed among treatment arms following 20 and 32 weeks of randomized therapy.

B. RATIONALE

Metformin and sulfonyl ureas, such as glyburide, are a known and effective combination in the treatment of type 2 diabetes mellitus. The two drugs have demonstrated a synergistic effect on glucose lowering when used in combination. Either drug can be used alone as first line monotherapy. They may also be used in combination with each other if monotherapy of either is inadequate. No data currently exists on the use of low dose combination therapy for first line use.

Treatment with a fixed dose combination tablet was expected to improve glycemic control as first line therapy in subjects with type 2 diabetes mellitus with inadequate control on diet and exercise. Glycemic control was expected to be achieved at lower doses than monotherapy with comparable or less potential side effects of the individual agents and with ease of administration.

This randomized double-blind, placebo-controlled study in subjects with type 2 diabetes mellitus who have inadequate glycemic control on diet and exercise tested the following hypotheses:

1. Administration of a fixed dose metformin/glyburide combination product for 20 weeks (4 weeks stable once daily dosing in Period B and 16 weeks of treatment in Period C) in subjects with type 2 diabetes mellitus who have inadequate glycemic control on diet and exercise will produce significant reductions in HbA_{1c} compared to placebo.
2. Administration of a fixed dose metformin/glyburide combination product for 32 weeks in subjects with type 2 diabetes mellitus who have inadequate glycemic control on diet and exercise will be well tolerated.

C. OBJECTIVES(1) Primary

To compare, after 20 weeks of oral administration, the effect of 2 dosage strengths (Examples 1 and 2) of a fixed combination metformin/glyburide tablet that has been titrated for glycemic control on the reduction in HbA_{1c} level versus placebo.

(2) Secondary (included the following)

1. To assess safety and tolerability among treatment arms after 20 and 32 weeks of randomized therapy. Glycemic control may be achieved with a similar incidence in hypoglycemia with the fixed dose combinations as compared with sulfonyl urea alone or decreased gastrointestinal side effects as compared with metformin alone.
2. To assess after 20 weeks and assess after 32 weeks, the proportion of subjects with a therapeutic response in glycemic control of oral administration of each metformin/glyburide combination regimen when compared to the therapeutic response achieved with metformin monotherapy, glyburide monotherapy and placebo regimens. Therapeutic plasma glucose response will be defined as a FPG < 126 mg/dL (based on current ADA guidelines for FPG). Therapeutic response for HbA_{1c} will be defined as HbA_{1c} < 7%.
3. To assess after 20 weeks and assess after 32 weeks, the reductions in fasting glucose and 2-hour postprandial glucose and insulin levels following the oral administration of each fixed combination metformin/glyburide regimen with the reduction in fasting glucose and 2-hour

postprandial glucose and insulin level achieved with metformin monotherapy, glyburide monotherapy and placebo.

- 5 4. To assess the durability of reductions in HbA_{1c} levels after 32 weeks of administration of fixed combination metformin/glyburide product.
- 10 5. To assess long-term safety and efficacy of fixed combination metformin/glyburide products.

D. STUDY DESIGN

This was a multicenter, randomized, five-arm, parallel group, double-blind, placebo controlled trial of the antihyperglycemic activity of a fixed combination metformin/glyburide tablet as first line therapy in subjects with type 2 diabetes mellitus who have inadequate glycemic control (HbA_{1c} < 7%), with diet and exercise. Patients were drug naive or had no oral antihyperglycemic therapy for the 2 months prior to screening. Approximately 100 US sites enrolled up to a maximum of 800 patients with type 2 diabetes mellitus who had inadequate glycemic control defined as an HbA_{1c} between 7-11% on diet and exercise. The minimum number of patients required to achieve the primary outcome was a total of 500 patients or 100 patients per arm. However, recruitment continued for up to 6 months to recruit up to a maximum of 150 patients per arm to provide additional safety information. The design included 3 periods as follows:

(1) Period A - Two Week Dietary and Placebo Lead-in Phase

This initial phase included dietary instruction on a eucaloric, weight maintaining diabetes prudent diet consistent with ADA guidelines or a balanced diet of approximately 55% carbohydrates, 20% protein and 25% fat.

Tolerability of the administration of multiple capsules and tablets were assessed with placebo. Home glucose meters were dispensed with instruction on their use.

5 **(2) Period B - 4 Week Double-blind Once Daily**
 Stable Dose Phase

 Period B began the randomized, double-blind, parallel quadruple dummy treatment phase. Eligible patients were randomized to 1 of 5 study arms which
10 included placebo, glyburide monotherapy, metformin monotherapy, and two different dose strengths of fixed combination metformin/glyburide product (Examples 1 and 2). Subjects were maintained on once daily dosing for a 4 week period so that the contribution of the individual
15 components of the combination product can be assessed by short term glycemic parameters.

 This 4 week phase at stable once daily dosing illustrated the contribution of the individual components of the fixed combination product using short term
20 glycemic parameters. Glycemic control was assessed with fructosamine and fasting glucose.

(3) Period C - 28 Week Double-blind Titration and
 Stable Dose Phase

25 Period C was the continuation of the randomized, double-blind treatment phase. Subjects were titrated for glycemic control over the first four weeks then dose was maintained for a 24 week stable dose treatment segment. Analysis for the primary outcome, the change from
30 baseline in HbA_{1c} of the two combination therapies (Examples 1 and 2) relative to placebo, was assessed at week 16 of Period C which was after 20 weeks of randomized, double-blind treatment. This was done at this time as there had been adequate time for
35 stabilization of HbA_{1c} and for safety reasons as it was anticipated that a high number of placebo treated patients may have had to discontinue randomized study

medication due to insufficient glycemic control as treatment duration was extended. Subjects not discontinuing randomized study drug due to lack of efficacy remained on stable doses for a total of 24 weeks to evaluate durability of efficacy and gather additional safety and tolerability data. The study remained blinded and those subjects who discontinued randomized study drug due to lack of efficacy were eligible to begin the long-term, open-label treatment phase with fixed combination product.

This 28 week phase included an initial 4 week titration segment to improve glycemic control followed by a 24 week stable dose phase. Analysis for the primary outcome was assessed at the 16th week of Period C. Subjects were evaluated for discontinuation of randomized study drug due to lack of glycemic control beginning at visit C1 through C85. Subjects were evaluated for lack of efficacy at visit C113 and all subsequent visits until the end of randomized treatment. The assignment of randomized study drug remained blinded. Subjects who remained on randomized study drug continued the stable dose phase for a total of 28 weeks to allow evaluation of durability of efficacy and to gather additional safety and tolerability data. Subjects were evaluated for discontinuation of study medication due to lack of glycemic control on or after Visit C1 (Week 0, Period C).

DOSING

Study drugs for this study were defined as: placebo, glyburide, metformin, metformin/glyburide 250/1.25 mg and metformin/glyburide 500/2.5 mg. For blinding purposes this study incorporated a quadruple-dummy design. Patients meeting the inclusion criterion without meeting any exclusion criterion, satisfying the Period A glycemic criteria, were eligible for enrollment into Period A.

Period A:

This period was a single-blind placebo lead-in to test patient tolerability of ingesting multiple capsules and tablets in addition to evaluating compliance with the quadruple dummy design. Patients received kits
5 containing four bottles of placebo corresponding to study drug.

Week 0 (Visit A1) - Subjects were instructed to take 1 capsule or tablet from each bottle with their
10 first meal of the morning.

Week 1 (Visit A8) - Subjects were instructed to take 1 capsule or tablet from each bottle with their first meal of the day and a second capsule or tablet from each bottle with their evening meal.
15

Period B:

Following completion of the single-blind lead-in phase (Period A), qualifying subjects commenced therapy in the randomized, double-blind treatment phase (Period
20 B). At visit A15/B1 subjects were randomized to once daily dosing with breakfast of placebo, glyburide 2.5 mg, metformin 500 mg, metformin/glyburide 250/1.25 mg or metformin/glyburide 500/2.5 mg. Once daily dosing remained stable for a total of 4 weeks.
25

Period C:

Following completion of the 4 week stable once daily dose phase (Period B) subjects continued the same randomized therapy in the 28 week titration/stable dose
30 treatment phase (Period C). Study medication was titrated at visits C1, C15 and C29. Medication was dosed with the first morning meal and with the evening meal. Potential maximal doses achieved included glyburide 10 mg, metformin 2000 mg, metformin/glyburide 1000/5 mg,
35 metformin/glyburide 2000/10 mg. After the 4 week titration segment in Period C, subjects continued on a

stable dose of study medication for the remainder of Period C.

Once adequate glycemic control had been achieved or maximum dose had been attained, study drug was not
5 increased and was only reduced with documented hypoglycemia.

RESULTS

The results obtained from the above studies
10 indicate that the low dose metformin-glyburide (250/1.25) formulation of the invention achieved glycemic control at least essentially equivalent to the high dose metformin-glyburide (500/2.5) formulation as evidenced by

15 (1) a therapeutic response for hemoglobin A1c, namely, a reduction in HbA1c of below 7% (from a mean baseline of 8.2%) at week 20 (Figures 1, 2 and 3), at weeks 20 and 32 and final visit (Figures 4 and 5)

(2) a therapeutic response for fasting plasma
20 glucose (FPG), namely, a reduction in FPG to less than 126 mg/dL after 20 weeks (from a baseline of about 175 mg/dL), (as shown in Figures 6)

(3) a therapeutic response for post-prandial
25 insulin levels, namely an increase in post-prandial insulin of 19-25 μ iu/mL (microinternational units/mL) (Figure 7)

(4) a therapeutic response for post-prandial
glucose excursion (PPG) (that is the difference between
30 post-prandial glucose and fast plasma glucose), namely, a decrease in post-prandial glucose excursion at week 20 of 17.7 for the 500/2.5 mg combo and 20.8 for the 250/1.25 mg combo versus 15.2 for metformin, 6.8 for glyburide. (Figures 8A and 8B).

At the same time, the above efficacy results employing the low dose formulation of the invention (Example 1) were achieved with reduced incidence of side effects (Figures 9 and 10).

5 As seen in Figure 9, the incidence of hypoglycemia employing the low dose formulation of the invention (Example 1) is less than about 1/3 of that occurring using the prior art high dose formulation (Example 2) employed in generally accepted medical practice for
10 treating diabetes.

 As seen in Figure 10, the incidence of gastrointestinal side effects employing the low dose formulation of the invention (Example 1) is less than 20% of that occurring using the high dose formulation
15 (Example 2) employed in generally accepted medical practice for treating diabetes.

 A discussion of the above results follows.

Discussion of Results

20 The progression to clinical type 2 diabetes takes time and requires the presence of multiple physiologic defects which are already present by the time most individuals are diagnosed with diabetes. Oral
therapeutic options for the treatment of type 2 diabetes,
25 until the last few years, have been severely limited. Further, with continued disease progression over time, all oral antihyperglycemic therapies are expected to become less effective leading to inadequate glycemic control for the patient.

30 Combination therapy has traditionally been indicated for second line use if initial single agent therapy is found to be ineffective, called "primary failure," or after initially effective agents are found to be ineffective at maintaining glucose control, called
35 "secondary failure." Switching from one monotherapy that is failing to an alternative monotherapy has not been proven to be effective in achieving glycemic control;

only the addition of a second agent with a different mechanism of action has been shown to achieve improved glycemic control. Given that a combination of insulin resistance and relative deficiency of insulin secretion
5 is the pathophysiological basis of type 2 diabetes, it is expected that combinations of agents offer greater therapeutic potential. Thus, both clinical experience and pathophysiologic evidence support the use of combination therapy earlier in the disease process.

10 While a fixed combination of metformin and glyburide is not a novel concept, and, as discussed above, different forms of it are available outside the U.S. for first and second line therapy, the use of combination therapy, low or moderate dose, as first line
15 treatment in drug naïve patients has never been studied in large controlled clinical trials. Treating to a near euglycemic target, an HbA_{1c} < 7% as recommended by the ADA, is the goal with any antihyperglycemic therapy. However, depending upon the duration of diabetes and the
20 progression of the disease, a single agent may not provide the efficacy necessary to bring even newly diagnosed patients to their target goal. The data presented in this summary provides evidence that a low dose fixed combination metformin/glyburide product is
25 safe and provides the efficient antihyperglycemic potency necessary to bring most drug naïve patients to the ADA's recommended glycemic target.

As first line therapy, a single formulation of fixed combination metformin/glyburide in ratio of a 200:1
30 metformin/glyburide was evaluated using two different dose strengths, a low dose (metformin/glyburide 250/1.25 mg) and a medium dose (metformin/glyburide 500/2.5 mg). The two dose strengths of fixed combination metformin/glyburide product were compared in a double-
35 blind study to placebo, glyburide monotherapy and metformin monotherapy. Mean final doses achieved in each treatment arm were approximately 5.3 mg of glyburide,

1307 mg of metformin, 557/2.78 mg of low dose (250/1.25 mg) metformin/glyburide fixed combination and 818/4.1 mg of medium dose (500/2.5 mg) fixed combination. When used as first line therapy, fixed combination

- 5 metformin/glyburide treatment achieved statistically significant improvement in glycemic control compared to metformin, glyburide or placebo. The interim open-label treatment data confirmed the clinical utility of fixed combination therapy in a more "glycemically diverse"
- 10 patient population and for a longer period of time.

Safety

- As first line therapy use, two dose strengths of metformin/glyburide were evaluated; a low-dose (250/1.25
- 15 mg) and a medium dose (500/2.5 mg) strength were compared with placebo, glyburide and metformin. In the double-blind phase of this study, diarrhea was the most frequently-occurring adverse effects (AE) in those subjects who were on metformin mono- or combination
- 20 therapy. Importantly, however, the incidence of gastrointestinal AEs was lower in the low dose fixed combination group than in the metformin monotherapy group (as seen in Figure 10). Discontinuations due to AEs also occurred with the lowest frequency in the low dose fixed
- 25 combination group compared to any of the other active treatments. Discontinuations due to lack of glycemic control were lowest in both the fixed combination groups, and severe hypoglycemia was not observed in this study. The frequency of subjects reporting an episode of
- 30 hypoglycemia was highest in the medium dose fixed combination treatment group, while the low dose group had a lower incidence than glyburide monotherapy (Figure 9). Mild increase in lactate levels were observed in all metformin groups, but no cases of lactic acidosis were
- 35 reported in this study.

In the open-label phase of the study, subjects could be directly enrolled if they did not meet the

glycemic criteria for entry into the double blind study. Subjects could also enter the open-label phase if they discontinued prematurely from the double-blind phase due to lack of glycemic control, or after they completed the double-blind phase. In the open-phase of the study, the AE profile was similar to that observed in the double-blind phase, with the most frequently-occurring AEs in the same body systems. The low dose combination group again displayed a favorable overall safety profile compared to the medium dose group.

In both the newly-diagnosed subjects as well as inadequately-controlled subjects, the overall pattern of safety and tolerability observed in the double-blind studies was as expected from the clinical experience with metformin and glyburide. No new or unexpected events or laboratory abnormalities were observed in this clinical program. Interim analyses of the long-term open-label extensions support the favorable safety profile observed in the short-term phase of the studies. In particular, the low dose fixed combination showed a favorable safety/tolerability profile when compared to the other regimens used in this program.

Efficacy

Double-blind, first line therapy demonstrated a statistically significant mean decrease in HemoglobinA_{1c} (HbA_{1c}) of 1.3% from placebo for both fixed combination treatment groups and a mean decrease from baseline of approximately 1.5%. While all active therapy treatment groups achieved acceptable glycemic control, greater mean decreases in HbA_{1c} for both fixed combination treatment groups were achieved when compared to metformin therapy of glyburide therapy. Antihyperglycemic durability was observed with all active treatment groups (glyburide, metformin, metformin/glyburide 250/1.25 mg, metformin/glyburide 500/2.5 mg) as evidenced by the maintenance of the mean HbA_{1c} levels from Week 20 (6.64%,

6.79%, 6.68%, 6.44%) to Week 32 (6.78%, 6.96%, 6.87%, 6.68%) of double-blind therapy below the therapeutic target of 7% (Figures 3 and 4).

Interim open-label first line therapy data demonstrate that for subjects directly enrolled, the mean HbA_{1c} at baseline was 10.6%, and for the subset of subjects with available data, a mean decrease of 3.5% in HbA_{1c} was achieved with a mean HbA_{1c} of 7.1% through 26 weeks. Of the subjects directly enrolled into open-label therapy, 87% received the medium dose 500/2.5 mg fixed combination as initial therapy and at the time of the interim report, the mean dose of fixed combination therapy was metformin/glyburide 1569/7.85 mg. For subjects with available open-label data completing the double-blind treatment phase and continuing into the open-label treatment phase, the mean HbA_{1c} at baseline was 8.32%. For all subjects reaching 13 weeks of therapy, a mean decrease of 1.76% in HbA_{1c} was achieved with the mean HbA_{1c} of 6.56%. Of the subjects completing the double-blind treatment phase and continuing into the open-label treatment phase, 78% received the low dose (250/1.25 mg) and 22% received the medium dose (500/2.5 mg) fixed combination as initial therapy. The mean dose of fixed combination therapy was metformin/glyburide 696/3.48 mg.

No clinically significant patterns of greater or reduced effect were apparent in any of the sub-populations (age, gender, race) with respect to response in HbA_{1c} from baseline in either double-blind trial with fixed combination metformin/glyburide as first line therapy.

This clinical program also assessed fasting plasma glucose as a parameters of short term glycemic control. FPG results in double-blind studies were consistent with the HbA_{1c} results. As first line therapy, statistically and clinically significant larger mean decreases in FPG for both fixed combination treatment groups compared to placebo and metformin were achieved (Figure 6). An early

response to fixed combination therapy was observed; differences among treatment groups were apparent by Week 2 of double-blind therapy as a time when subjects were still undergoing initial titration and were receiving only one-half potential maximum dosing. This early response at one-half maximum dosing in a monotherapy refractory patient population demonstrates the benefit of combination therapy for the patient and using combination therapy earlier in the disease process.

HemoglobinA_{1c} is the prevailing standard measure of overall glycemic control and it is the glycemic marker found to be correlated with long term complications. Although, fasting plasma glucose, the current standard for the diagnosis of diabetes, is a faster, more convenient marker, it does not provide an optimal assessment of circadian glycemic control. It has been shown, and intuitively understood, that non-fasting plasma glucose is a better marker of diabetic control than FPG in type 2 diabetes; it also correlates better with HbA_{1c}. Postprandial hyperglycemia is an early marker of the metabolic defects found in type 2 diabetes and contributes to beta cell dysfunction. An important association between postprandial glucose levels and cardiovascular disease has been demonstrated. If normal glycemia is the goal in preventing long term complications of diabetes then monitoring and lowering postprandial glucose is a rational strategy in improving metabolic function and achieving overall glucose control.

As first line therapy, statistically significant larger mean decreases in absolute postprandial glucose (63-65 mg/dL) were observed for both fixed combination treatment groups than the placebo group. Larger mean decreases in absolute PPG were also achieved compared with glyburide (16-18 mg/dL) and metformin (18-20 mg/dL) monotherapy (Figures 8A and 8B). The 2-hour postprandial glucose excursion from a fasting baseline for both the low dose (22.5 mg/dL) and medium dose (23.9 mg/dL) fixed

combination treatment groups was only 56%-59% of placebo (40.3 mg/dL), 59%-63% of glyburide (38.2 mg/dL) and 75%-81% of metformin (29.5 mg/dL). Evaluating the excursion rather than the absolute value demonstrates that

5 glyburide is similar to placebo, metformin achieves better postprandial glucose lowering than glyburide and placebo, and that the low dose combination is the most powerful in lowering postprandial glucose excursion. As there is no published clinical data with combination
10 therapy studied in a drug naive patient population, these results add new insight to understanding of the impact of treatment options at this stage of the disease. Indeed, the results could not have been predicted from the changes observed in the much studies second line therapy
15 population.

Insulin levels were evaluated in the fasting and postprandial state in the first line therapy study (Figure 7). There was a statistically significant increase in insulin response in the presence of a glucose
20 load for both fixed combination treatment groups (24-28.8 μ iu/mL) compared to placebo. A larger increase in insulin response in the presence of a glucose load for the low dose fixed combination (14.6 μ iu/mL) treatment group was observed when compared to glyburide monotherapy and a larger increase in insulin response in the presence
25 of a glucose load for both fixed combination (21-25.8 μ iu/mL) treatment groups was observed when compared to metformin monotherapy. When considering the mean doses of active therapy per treatment group, the insulin
30 response cannot be explained by the sulfonylurea component alone with fixed combination therapy. This clinical data supports preclinical work with isolated pancreatic islet cells where it has been suggested that metformin prevents the hyperglycemic desensitization of
35 the islet cells. The combination of the physiologic and appropriate increased insulin response with a corresponding larger decrease in glucose excursion

suggests that the combination is improving the efficiency of the pancreas in responding to a glucose load, preserving beta cell function and improving insulin sensitivity.

5 The essential goal in the management of patients with type 2 diabetes, in addition to aggressively treating elevated blood pressure and lipid levels, is achieving as near normal glycemic levels as possible or achieving glycemic therapeutic targets. There was a
10 greater response to fixed combination therapy with respect to greater frequencies of subjects achieving therapeutic targets and greater decreases in absolute HbA_{1c}. As first line therapy, a higher frequency of subjects on fixed combination therapy (66%-71%) achieved
15 a glycemic target of an HbA_{1c} \leq 7% compared with 60% of sulfonylurea monotherapy, 50% of metformin monotherapy and 20% of placebo following 20 weeks of double-blind therapy. Approximately 28% of subjects in each fixed combination group had decreases in HbA_{1c} from baseline
20 greater than 2.0%, compared with 16%-17% of each monotherapy group and 3% of placebo. Of note, is that these targets were not achieved with simply higher total doses of medication, but with lower doses of the complementary components. Mean final doses achieved in
25 each first line therapy treatment arm were approximately glyburide 5.3 mg, metformin 1307 mg, low dose fixed combination 557/2.78 mg and medium dose fixed combination 818/4.1 mg. For the change in HbA_{1c} by number of tablets, the pattern observed with fixed combination therapy is
30 not unexpected from a pathophysiologic viewpoint. It indicates that there is a clear response to target at all dose levels and that the need for higher doses correlates with a higher baseline HbA_{1c}. A similar pattern can be detected for glyburide up to a total dose 7.5 mg; no
35 clear pattern was observed with metformin therapy.

The data presented supports low dose fixed combination metformin/glyburide as the first line agent

most likely to bring a patient to therapeutic target, no matter how high their baseline HbA_{1c}. For both fixed combination therapies, the mean decrease from baseline HbA_{1c} is larger for subjects with higher baseline levels.

5 This phenomenon was not observed with glyburide, metformin or placebo and is not expected to be seen with other monotherapies. This demonstrates the contribution of components necessary for achieving therapeutic glycemic targets when baseline HbA_{1c} level is greater than
10 9%. Monotherapy was shown to have a plateauing of glycemic response for baseline HbA_{1c} levels < 9% while fixed combination therapy had additional incremental decreases in HbA_{1c} for baseline HbA_{1c} levels < 9%.

For all subjects enrolled into the open-label first
15 line treatment phase with available data for at least two time points, the mean HbA_{1c} at baseline was 9.45%. By Weeks 13, 26 and 39 approximately 50-55% of subjects had achieved an HbA_{1c} of less than 7% and an additional 30% had achieved an HbA_{1c} < 8%. This response rate and
20 magnitude of change in HbA_{1c} lowering can be expected with combination therapy but is rarely seen with monotherapy antihyperglycemic agents. The fundamental issue is what initial antihyperglycemic treatment will achieve the glycemic target of an HbA_{1c} < 7% in the greatest
25 proportion of patients. This data strengthens the need for the reevaluation of current type 2 diabetes treatment paradigms and to shift to the use of combination therapy sooner in the disease process. .

Weight gain is typically observed with all
30 antihyperglycemic agents other than metformin monotherapy. With improved glycemic control, a weight gain is actually expected as calories are conserved rather than lost due to poor metabolic control. In this clinical program, as glycemic control improved, minimal
35 early weight gain of approximately 1-2 kg was observed with fixed combination therapy; this was comparable to the 2 kg weight gain observed with first line glyburide

monotherapy. In double-blind therapy, after the initial minimal gain, weight remained stable and did not continue to increase with time.

Overall there were no clinically or statistically
5 significant differences between any of the treatment groups with respect to changes in the plasma lipid profile. As the most severe patients were excluded from the placebo controlled trial, smaller changes in response to therapy might be undetectable. The first line therapy
10 patient population had inadequate glycemic control but diet and exercise has already succeeded in bringing the mean HbA_{1c} to 8.2%. In subjects treated with fixed combination therapy, there was no adverse effect on the plasma lipid profile (total cholesterol, LDL, HDL, and
15 triglycerides) or significant differences compared with placebo or either glyburide and metformin monotherapy.

With better understanding of the relationship between diabetes control and long-term complication rate the goal of diabetes management today is to achieve and
20 maintain as near normal glycemia as possible. Targeting multiple defects using agents with synergistic or complementary mechanisms of action intuitively makes sense to achieve a therapeutic glycemic target. Improved understanding of the natural history of type 2 diabetes
25 suggests that current treatment paradigms of allowing "failure" to occur prior to implementing a more aggressive treatment strategy should be reassessed. Earlier use of low dose combination therapy, particularly when the use of lower doses results in better
30 tolerability, therefore appears to be an important therapeutic approach if targets are to be achieved and compliance maintained. The fixed combination evaluated in this study allows for lower dosing and the ease of use in a single entity.

35 Low dose fixed combination metformin/glyburide therapy is safe and effective in achieving and maintaining glycemic control in patients with type 2

diabetes who have inadequate glycemic control with diet and exercise. The use of combination therapy earlier in the diabetes disease progression appears to be a clinically sound alternative to the classic treatment paradigms of allowing failure of step wise therapy before instituting a more aggressive, but clinically sound, treatment strategy. Though not evaluated in this short-term study, the strategy to achieve as near normal glycemic targets as possible is likely to have an impact in slowing the progression of the diabetes disease process and delay the onset of long-term diabetes complications. Given a refractory monotherapy patient population the fixed combination of metformin and glyburide was associated with a clinically significant improvement in glycemic control without evidence of detrimental metabolic effects or safety concerns. There was no clinically significant hypoglycemia, no negative impact in plasma lipids and a limited early weight gain followed by stable weight with time. The synergism of the metformin and sulfonylurea combination is an established one; a fixed combination of metformin and glyburide is effective in improving glycemic control and is a rationale choice in the antihyperglycemic armamentarium. It is assumed that a fixed combination simplifies dosing, is more convenient and therefore may lead to better compliance with therapy.

The low dose (250/1.25 mg) fixed combination would be the initial starting dose as first line therapy in drug naive subjects. This should then be titrated as indicated to achieve a $HbA_{1c} < 7\%$.

OVERALL CONCLUSIONS

The safety and efficacy data presented from this clinical program assessing fixed combination metformin/glyburide as first line therapy in patients with type 2 diabetes confirm the following:

- The percentages of subjects who discontinue from therapy because of hyperglycemia were lower for fixed combination metformin/glyburide compared with metformin, glyburide, and placebo.
- 5
- Hypoglycemia and symptoms of hypoglycemia, as first line therapy (Figure 9), occurred less often with metformin/glyburide 250/1.25 mg compared to metformin/glyburide 500/2.5 mg and glyburide.
- 10
- As first line therapy, the incidence of gastrointestinal adverse events associated with fixed combination was lowest for metformin/glyburide 250/1.25 mg compared with metformin/glyburide 500/2.5 mg and metformin (Figure 10).
- 15
- No new or unexpected adverse events or laboratory abnormalities occurred in subjects who received long-term open-label fixed combination metformin/glyburide.
- 20
- Significantly better efficacy of fixed combination metformin/glyburide at any dose strength as evidenced by greater reductions of all glycemic parameters (HbA_{1c}, postprandial glucose, fasting glucose and fructosamine) compared to placebo, glyburide and metformin therapy.
- 25
- A synergistic effect of the low dose combination in targeting multiple metabolic defects to improve beta cell function and insulin sensitivity, as evidenced by postprandial plasma glucose and insulin excursions, to achieve improved metabolic function and glycemic control.
- 30
- A higher frequency of patients on fixed combination metformin/glyburide therapy achieved a glycemic therapeutic target of an HbA_{1c} ≤ 7%.
- 35

- Efficient glycemic lowering to therapeutic targets for any baseline HbA_{1c} compared with placebo, glyburide and metformin therapy. As initial therapy, glyburide and metformin were shown to have a plateauing of glycemic response for baseline HbA_{1c} levels > 9% while fixed combination metformin/glyburide therapy had additional incremental decreases in HbA_{1c} for baseline HbA_{1c} levels > 9%.
- Limited early weight gain paralleling improved glycemic control, comparable to glyburide monotherapy; however, weight remained stable with time.
- No adverse effect of the fixed combination therapies on the lipid profile (total cholesterol, LDL, HDL, and triglycerides) or significant differences from placebo or either glyburide and metformin monotherapy.
- The favorable efficacy and tolerability of fixed combination metformin/glyburide 250/1.25 mg supports its use as the initial starting dose in first line therapy.

The above results clearly show that treating diabetes with the low dose metformin/glyburide formulation of the invention (250 mg/1.25 mg) is at least equivalent in efficacy to the higher dosage form (500 mg/2.5 mg), while resulting in reduced side effects.

What is Claimed is:

1. A method for first line treatment of type 2 diabetes, in a drug naive human patient, which comprises administering to a drug naive human patient in need of
5 treatment, as first line therapy, a low dose of a combination of metformin and glyburide.

2. The method as defined in Claim 1 wherein the daily dosage of metformin administered is less than 800
10 mg.

3. The method as defined in Claim 1 wherein the low dose combination of metformin and glyburide provides at least substantially equivalent efficacy in treating
15 diabetes in drug naive patients, but with substantially reduced side effects, as do combinations of metformin and said glyburide employed in substantially higher daily dosages as prescribed in generally accepted medical practice for first line therapy in treating diabetes.

4. The method as defined in Claim 1 wherein the starting daily dosage of metformin is as low as about one-fifth of the starting daily dosage of metformin employed in generally accepted medical practice for first
25 line therapy for treating diabetes.

5. The method as defined in Claim 4 wherein the daily maintenance dosage of metformin employed is up to that employed in generally accepted medical practice for
30 first line or second line therapy for treating diabetes.

6. The method as defined in Claim 1 wherein the starting daily dosage of glyburide is as low as about one-fifth of the starting daily dosage of glyburide
35 employed in generally accepted medical practice for first line therapy for treating diabetes.

7. The method as defined in Claim 6 wherein the daily maintenance dosage of glyburide employed is up to that employed in generally accepted medical practice for first line therapy or second line therapy for treating
5 diabetes.

8. The method as defined in Claim 1 wherein the metformin is administered in a daily dosage in an amount within the range from about 160 mg to about 750 mg, and
10 the glyburide is administered in a daily dosage in an amount within the range from about 0.5 to about 15 mg.

9. The method as defined in Claim 1 wherein the low dose combination of metformin and glyburide is
15 formulated as a single dosage form.

10. The method as defined in Claim 1 wherein the metformin is employed in a weight ratio to glyburide within the range from about 400:1 to about 50:1.
20

11. The method as defined in Claim 1 wherein the metformin and glyburide are employed in a weight ratio to each other of about 200:1 or 100:1.

12. The method as defined in Claim 1 wherein the metformin is administered in an amount within the range from about 125 to about 750 mg, one to four times daily, provided that the maximum daily dosage for metformin is about 750 mg per day, but more than about 225 mg, and the
25 glyburide is administered in an amount within the range from about 0.75 to about 5 mg, one to four times daily, up to a maximum of 15 mg per day.
30

13. The method as defined in Claim 1 wherein the metformin is administered in an amount within the range
35 from about 250 to about 500 mg, and the glyburide is

administered in an amount within the range from about 1.25 to about 5 mg.

14. The method as defined in Claim 1 wherein the
5 metformin/glyburide combination contains of 250 mg
metformin/1.25 mg glyburide.

15. The method as defined in Claim 1 wherein the
metformin/glyburide combination contains 500 mg
10 metformin/2.5 mg glyburide.

16. The method as defined in Claim 1 wherein the
metformin/glyburide combination contains 500 mg
metformin/5 mg glyburide.
15

17. The method as defined in Claim 1 wherein a
metformin/glyburide 250/ mg/1.25 mg dosage is
administered once a day or twice a day.

20 18. The method as defined in Claim 19 wherein the
metformin/glyburide 250 mg/1.25 mg dosage is administered
to patients with a baseline HbA_{1c} > 9% or a fasting
glucose > 200 mg/dL twice daily, with dosage increases,
where necessary, in increments of 250 mg/1.25 mg every 2
25 weeks, up to the minimum effective daily dose necessary
to achieve adequate glycemic control.

19. The method as defined in Claim 1 wherein said
metformin is employed in a daily dose as employed in
30 generally accepted medical practice for first line
therapy or second line therapy for treating diabetes.

20. The method as defined in Claim 1 wherein
glyburide is employed in a daily dose as employed in
35 generally accepted medical practice for first line
therapy or second line therapy for treating diabetes.

21. The method as defined in Claim 1 wherein the low dose combination comprising 250 mg metformin and 1.25 mg glyburide characterized in that it has at least substantially equivalent efficacy to a formulation
5 comprising 500 mg metformin and 2.5 mg glyburide in treating diabetes with respect to decrease in hemoglobin A_{1c}, decrease in insulin resistance, increase in post-prandial insulin levels and/or decrease in post-prandial glucose excursion, while providing substantially reduced
10 incidence of adverse side effects which are hypoglycemia and gastrointestinal side effects.

22. A method for first line treatment of type 2 diabetes, in a drug naive human patient, which comprises
15 administering to a drug naive human patient in need of treatment, as first line therapy, a low dose of a combination of metformin and glyburide wherein the starting daily dosage is 250 mg metformin and 1.25 mg glyburide.
20

23. A method for first line treatment of type 2 diabetes, in a drug naive human patient, which comprises administering to a drug naive human patient in need of treatment, as first line therapy, a therapeutically
25 effective low dose of a combination of metformin and glyburide wherein the starting daily dosage is 250 mg metformin and 1.25 mg glyburide twice a day or 500 mg metformin and 2.5 mg glyburide once a day.

30 24. A method for first line treatment of type 2 diabetes, in a drug naive human patient, which comprises administering to a drug naive human patient in need of treatment, as first line therapy, a therapeutically effective low dose of a combination of metformin and
35 glyburide wherein the starting daily dosage is 500 mg metformin and 5 mg glyburide.

25. A method for first line treatment of type 2 diabetes, in a drug naive human patient, which comprises administering to a drug naive human patient in need of treatment, as first line therapy, a therapeutically effective low dose of a combination of metformin and glyburide wherein the glyburide is such that the glyburide bioavailability is comparable to the glyburide bioavailability obtained with a separate administration of metformin and glyburide.

26. The method as defined in Claim 25 where in the combination of metformin and glyburide the particle size distribution of the glyburide is such that at most 10% of the particles are less than 2 μm and at most 10% of the particles are greater than 60 μm .

27. The method as defined in Claim 25 wherein the glyburide has a particle size distribution such that at most 10% are less than 3 μm and at most 10% are greater than 40 μm .

28. The method as defined in Claim 25 wherein the glyburide has a particle size distribution such that at most 25% are less than 11 μm and at most 25% are greater than 46 μm .

29. The method as defined in Claim 25 wherein 50% of the glyburide patients are less than 23 μm .

30. The method as defined in Claim 25 wherein the glyburide has a particle size distribution of about 25% undersize value not more than 6 μm , about 50% undersize value 7 to 10 μm and about 75% undersize value not more than 23 μm .

31. The method as defined in Claim 25 wherein the starting daily dosage is 250 mg metformin/1.25 mg glyburide or 500 mg metformin/2.5 mg glyburide.

32. The method as defined in Claim 3 wherein the substantially reduced side effects are hypoglycemia and/or gastrointestinal side effects which are diarrhea, nausea/vomiting and/or abdominal pain.

33. The method as defined in Claim 32 wherein the incidence of hypoglycemia in drug naive patients resulting from use of the low dose metformin-glyburide combination is 1/3 or less than in patients treated with double the metformin-glyburide present in the low dose metformin-glyburide.

34. The method as defined in Claim 32 wherein the incidence of gastrointestinal side effects in drug naive patients resulting from use of the low dose metformin-glyburide combination is 20% less than in patients treated with twice the amount of each of the metformin-glyburide present in the low dose metformin-glyburide.

35. A method for lowering blood glucose in a hyperglycemic human patient, which comprises administering to a drug naive human patient in need of treatment, as first line therapy, a therapeutically effective amount of a low dose pharmaceutical formulation as defined in Claim 1.

36. A method for decreasing insulin resistance and/or decreasing hemoglobinA_{1c} and/or increasing post-prandial insulin levels and/or decreasing post-prandial glucose excursion, in a human patient, which comprises administering to a drug naive human patient in need of treatment as first line therapy a pharmaceutical formulation as defined in Claim 1.

METHOD FOR TREATING DIABETESAbstract of the Disclosure

5 A method is provided for first line treatment of
type 2 diabetes employing a combination of metformin and
glyburide. A method for treating diabetes in drug naive
human patients is also provided employing the above
formulation to reduce insulin resistance and/or post-
prandial glucose excursion and/or hemoglobin 1Ac, and/or
10 increase post-prandial insulin, thereby treating the
diabetes.

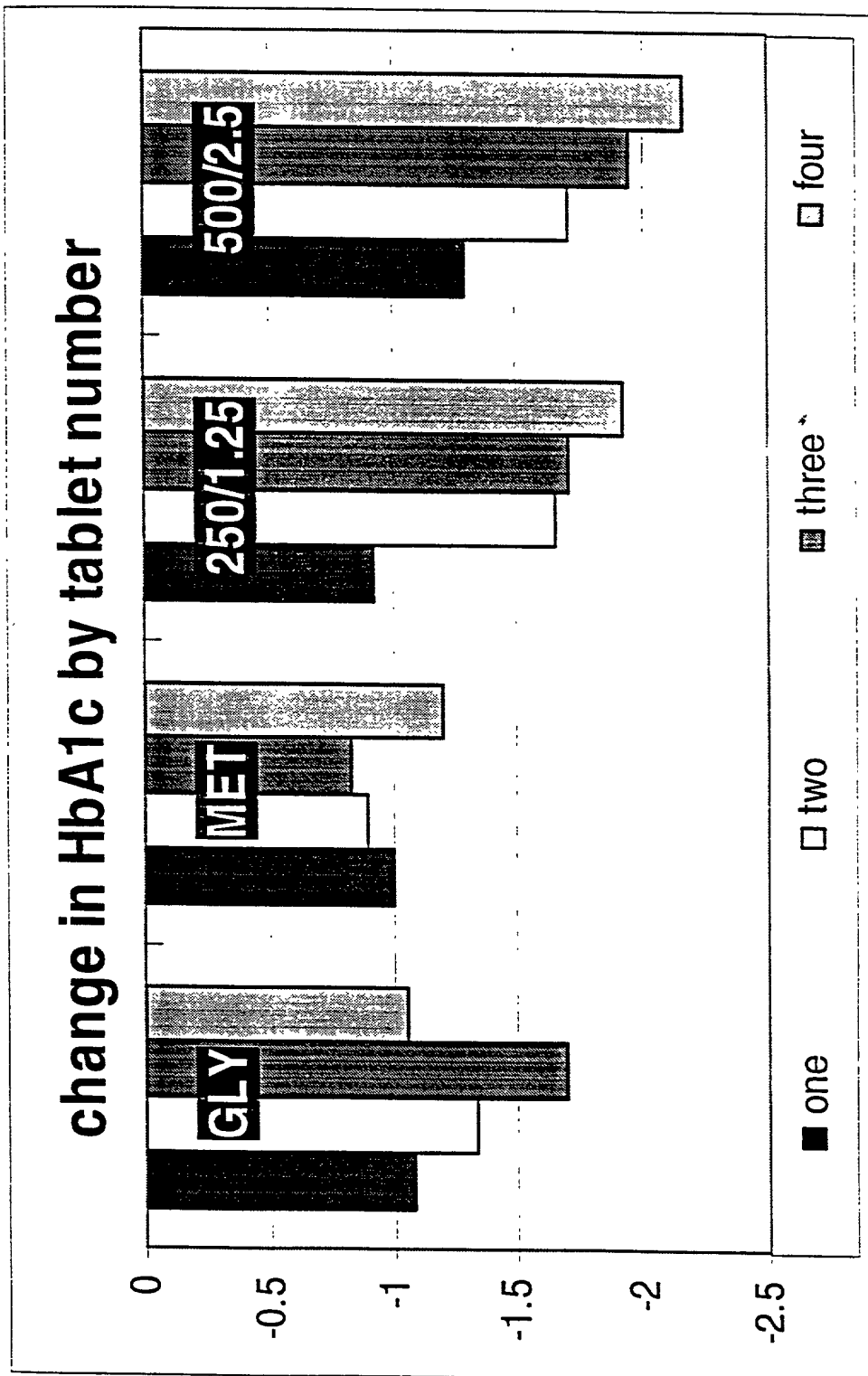


Fig 1

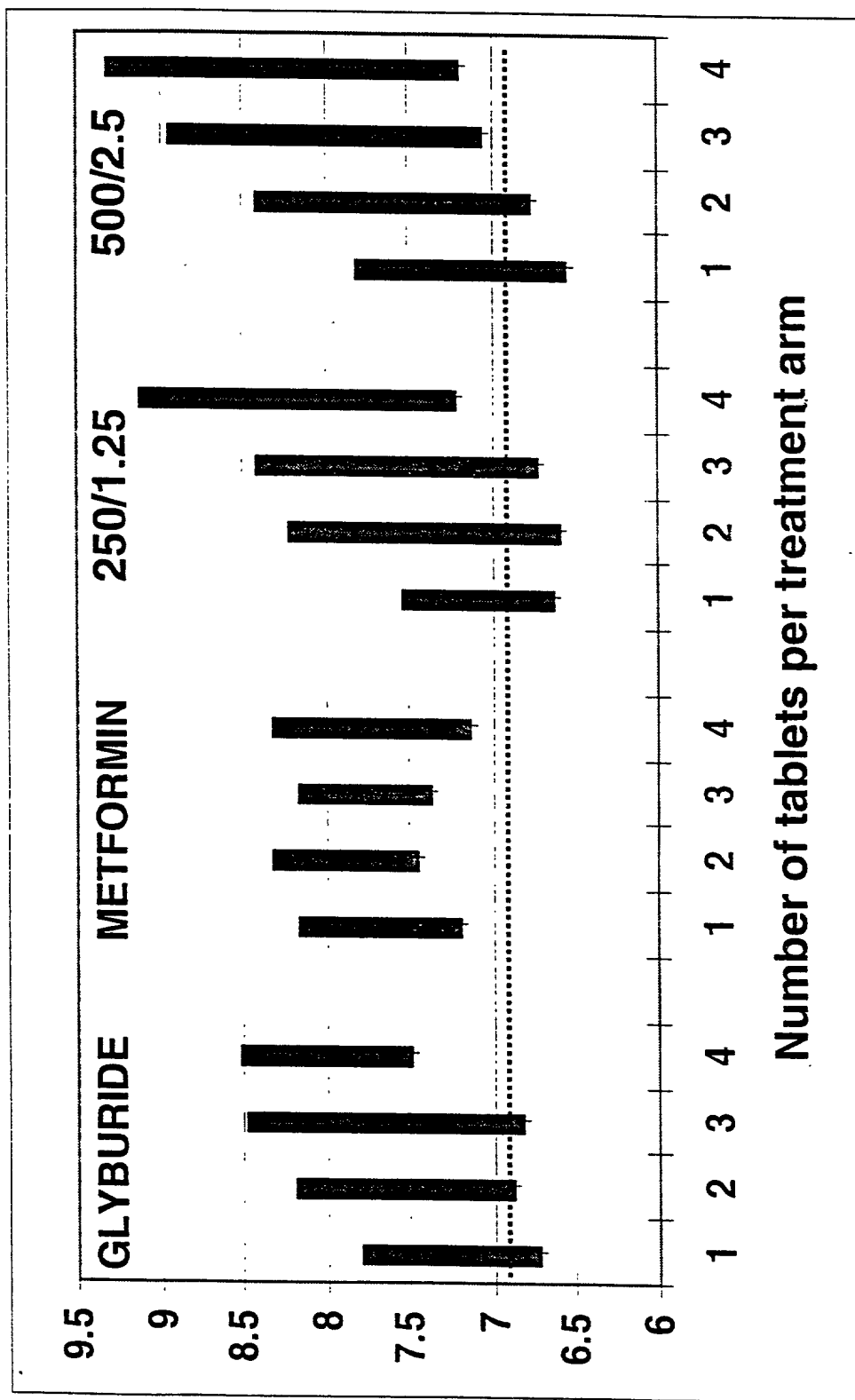


Fig 2

HbA1c over time

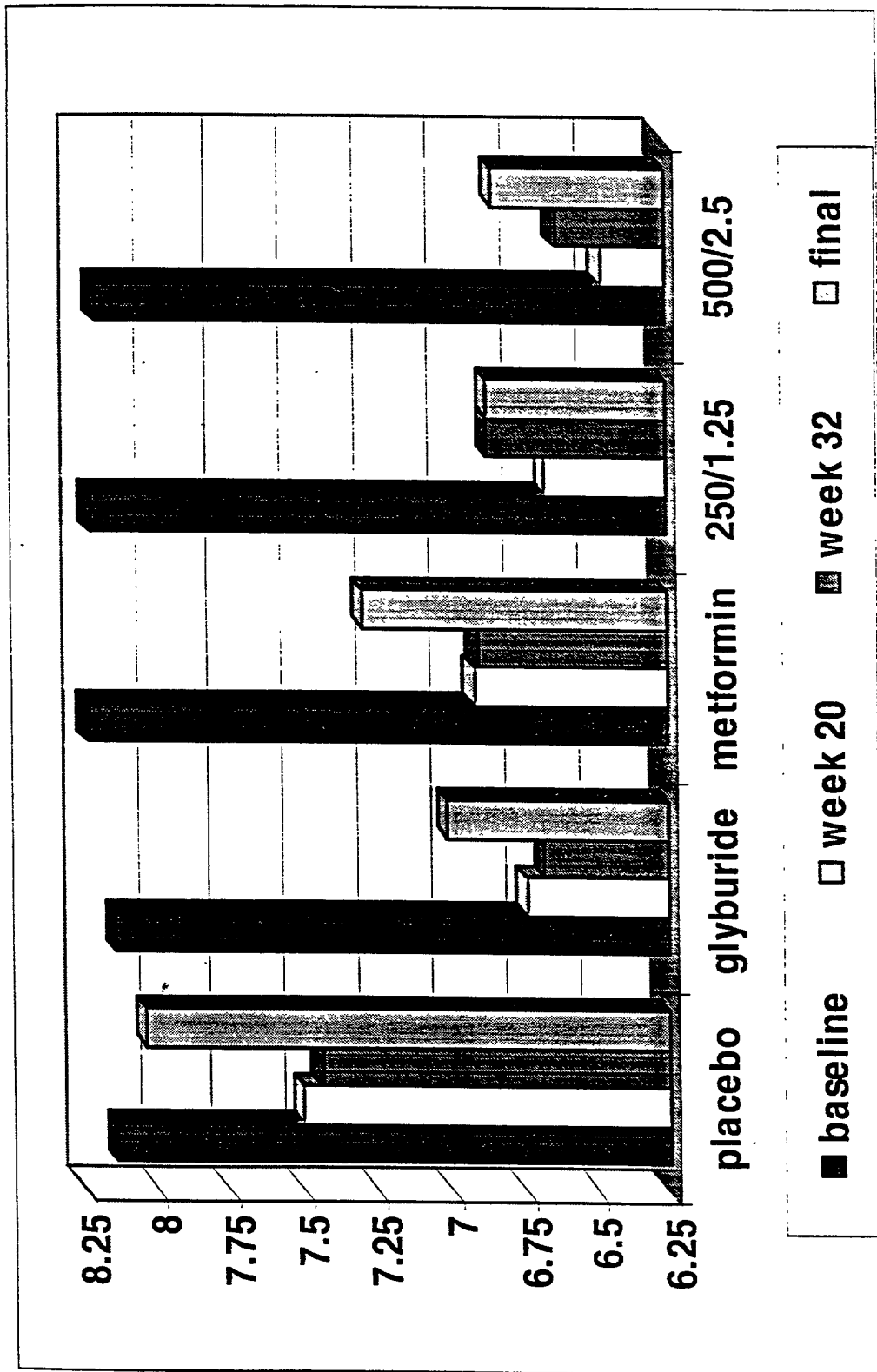


Fig 3

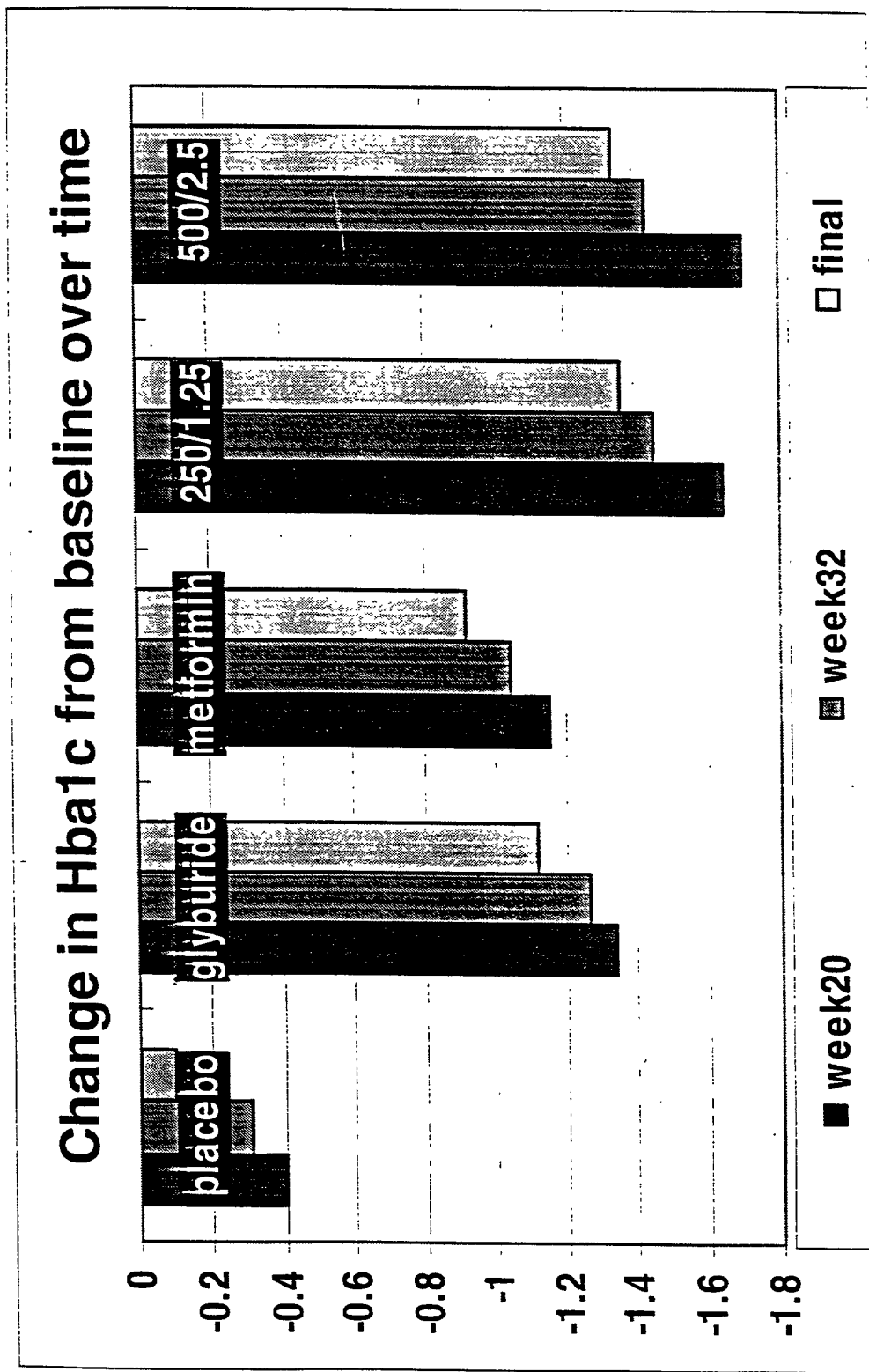


Fig 4

Change in HbA1c by baseline HbA1c

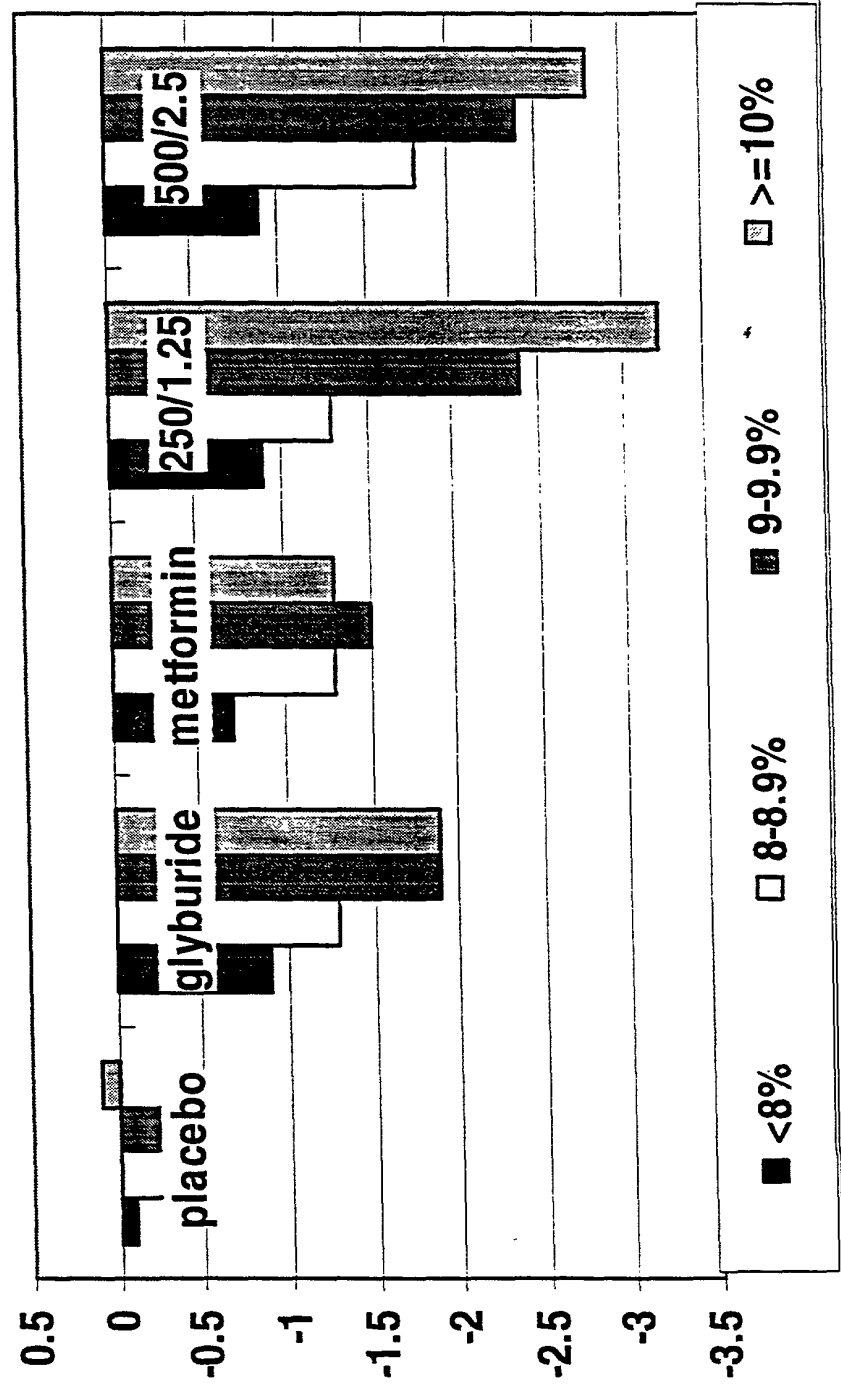


Fig 5

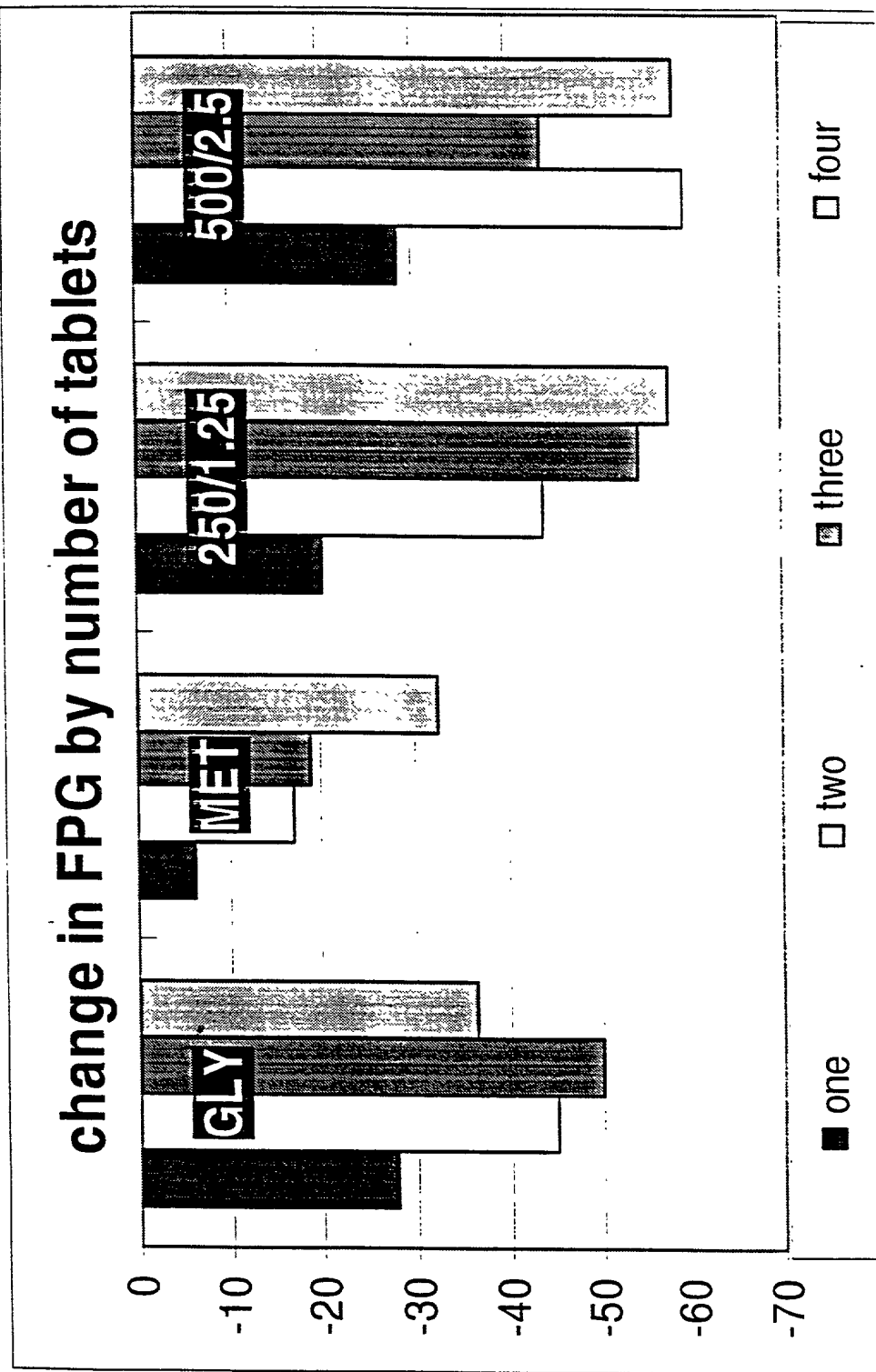


Fig 6

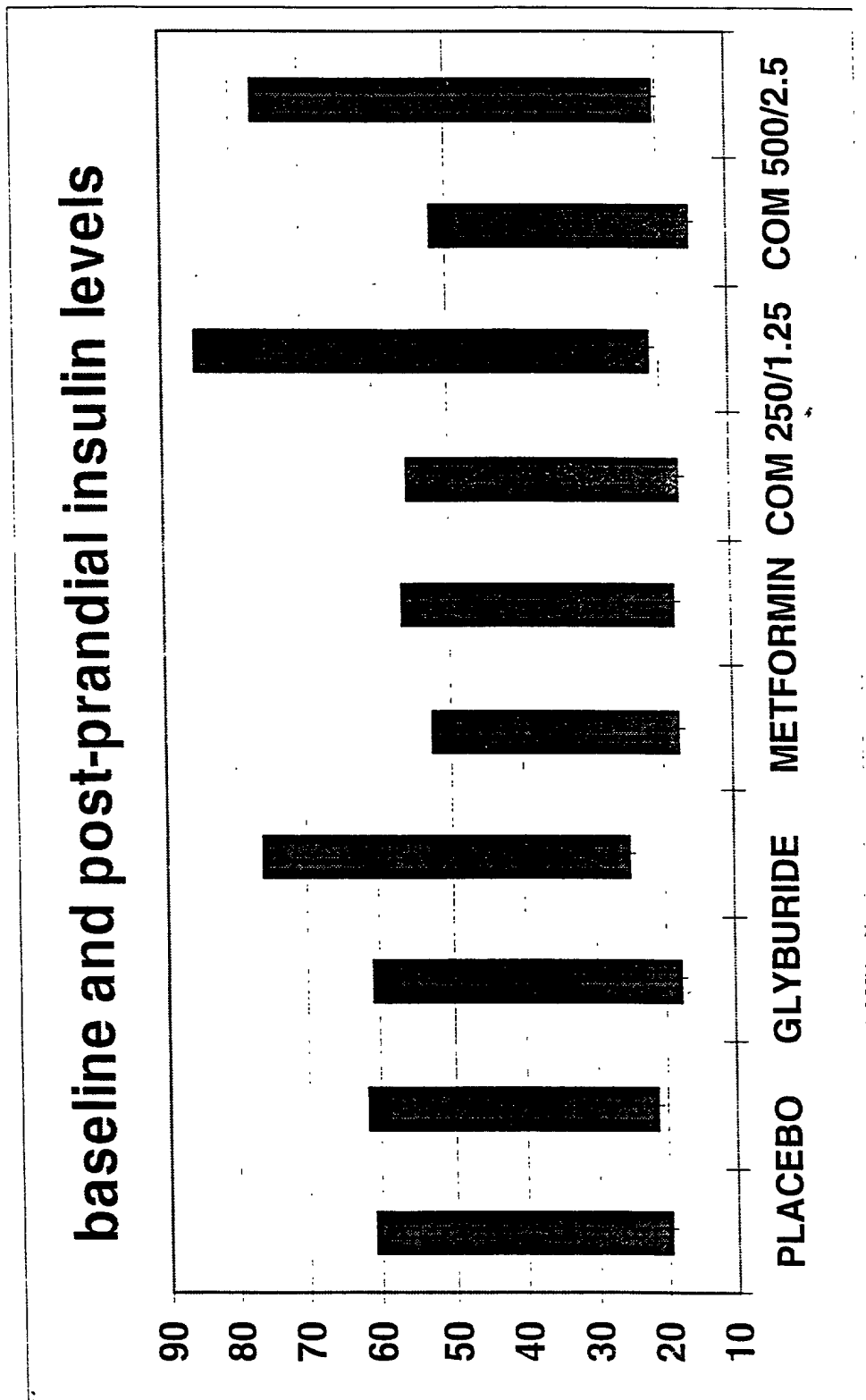


Fig 7

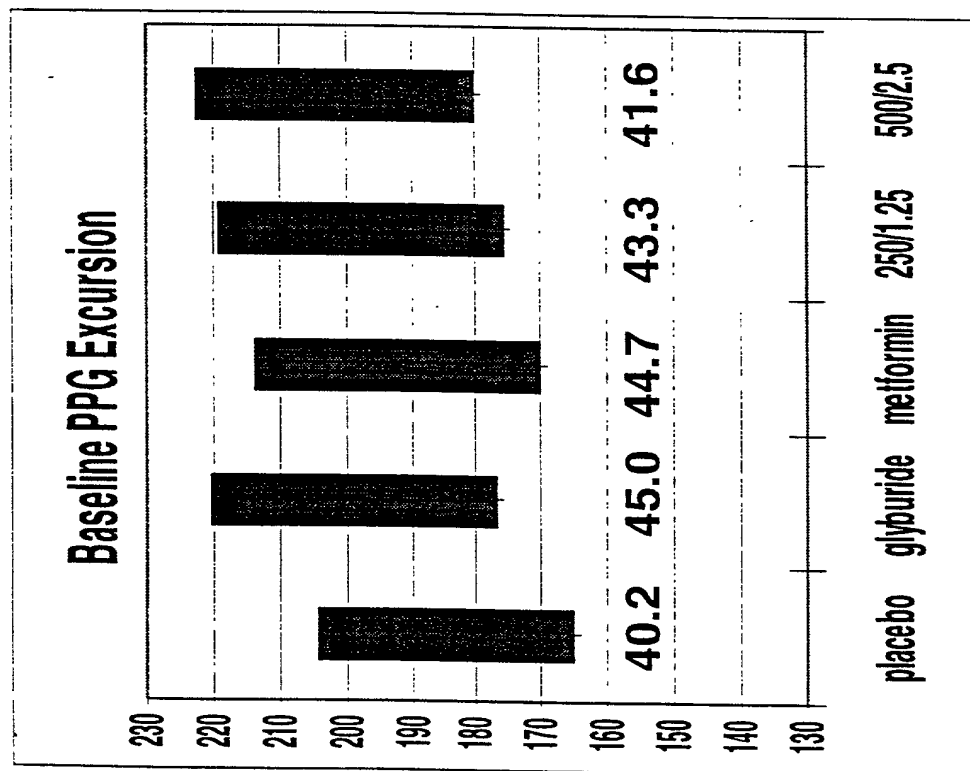


Fig 8A

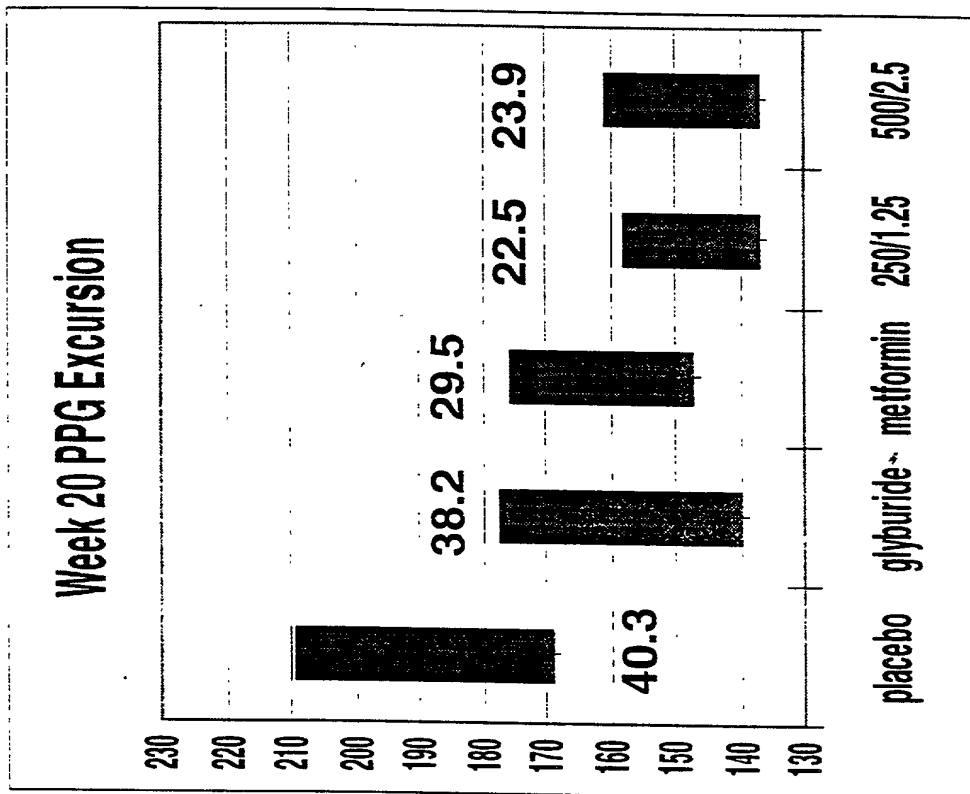
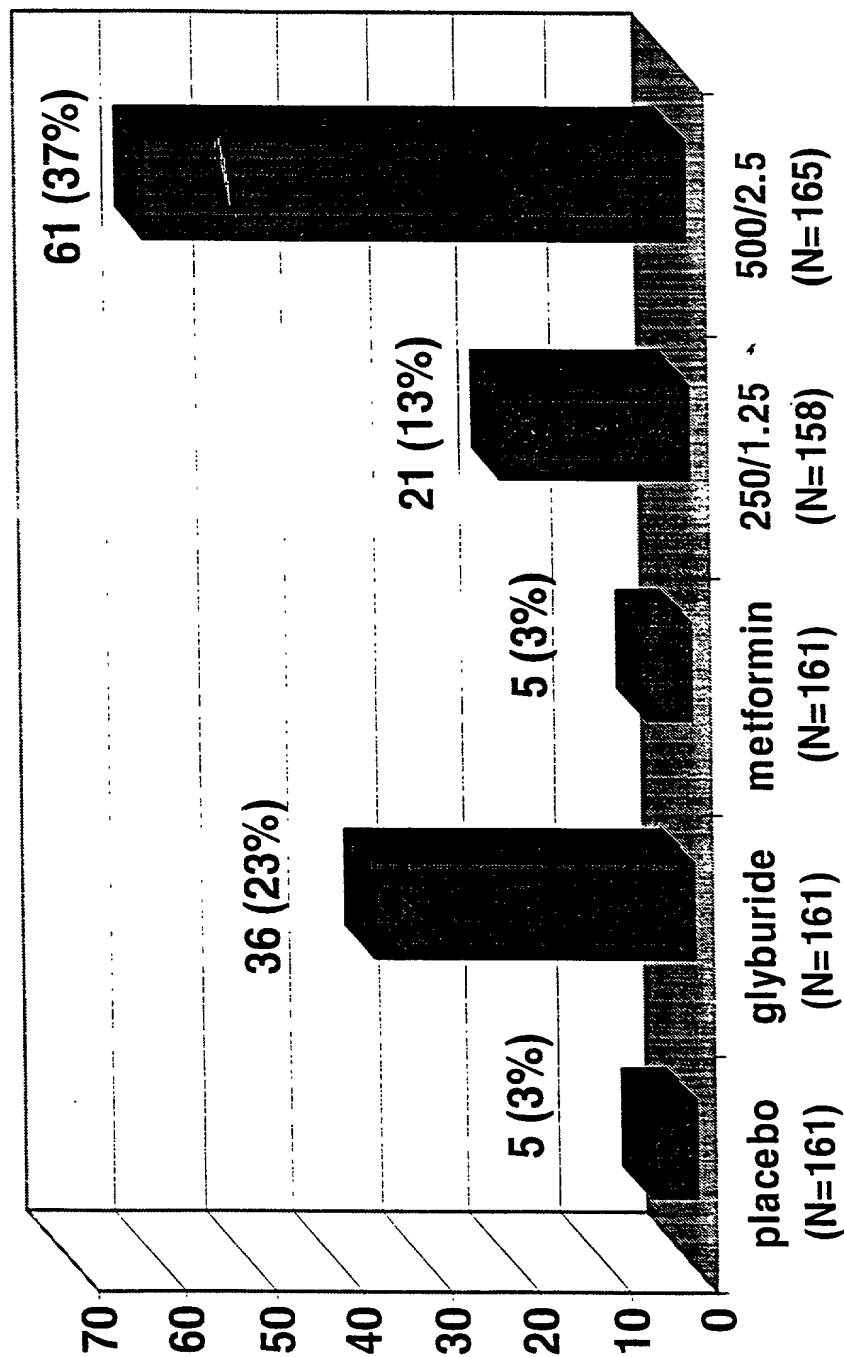


Fig 8B

Subjects reporting hypoglycemic symptoms



F159

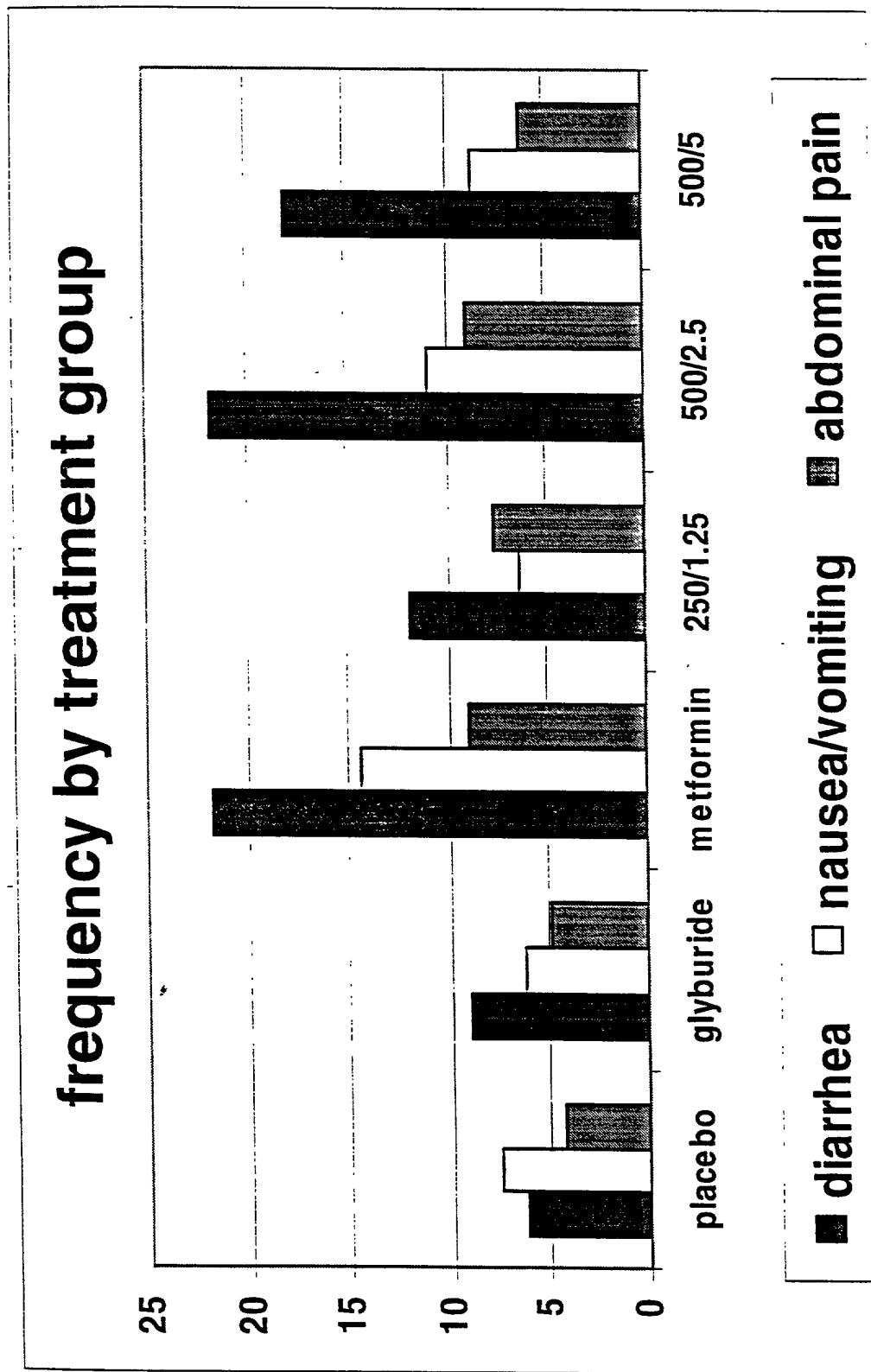


FIG 10

DECLARATION
AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: METHOD FOR TREATING DIABETES, the specification of which

X is attached hereto; or

 was filed on as U.S. Patent Application Serial No. / / .

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims.

I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to me to be material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119(a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

PRIORITY FOREIGN APPLICATION(S)
UNDER 35 U.S.C. §119(a)-(d)

<u>Number</u>	<u>Country</u>	<u>Filed</u> <u>(Day/month/year)</u>	<u>Priority</u> <u>Claimed</u> <u>(Yes or No)</u>
NONE			

I hereby claim the benefit under Title 35, United States Code, Section 119(e) of any United States provisional application(s) listed below:

PRIORITY U.S. PROVISIONAL APPLICATION(S)
UNDER 35 U.S.C. §119(e)

<u>Provisional Application No.</u>	<u>Filing Date</u>
NONE	

Continued on page 2

I hereby claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Section 112, I acknowledge the duty to disclose to the U.S. Patent and Trademark Office all information known to me to be material to the patentability of this application as defined in Title 37, Code of Federal Regulations, Section 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

PRIORITY U.S. APPLICATION(S)
UNDER 35 U.S.C. §120

<u>Application Serial No.</u>	<u>Filing Date</u>	<u>Status (patented, pending or abandoned)</u>
09/432,465	11/03/99	Pending

I hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Continued on page 3

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